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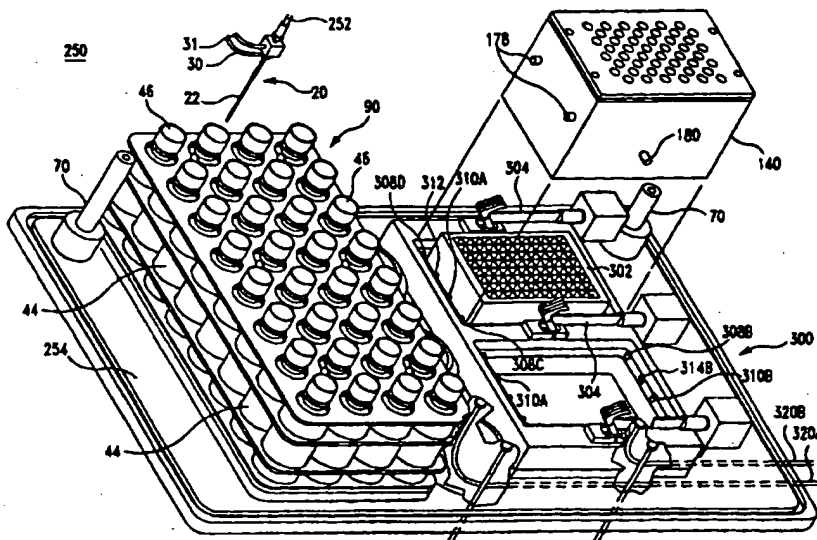
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## (57) Abstract

Method and apparatus for generation of chemical libraries are described. The apparatus includes a reaction block (140) which supports replaceable reaction chambers. Each reaction block is fitted with four sets of twelve reaction chambers. Reaction chambers are provided with s-shaped trap tubes which run into drain tubes. The reaction block is provided with gas lines and a septum seal such that gas pressurization empties the reaction chambers into the drain tubes. The drain tubes are arranged so as to mate directly with a standard 96 well microtiter plate for the collection of material. A docking station (300) secures registration of reaction blocks and introduces gases and liquids into the reaction blocks (140). An inert atmosphere is maintained in the reaction block by a top and an optional bottom seals. A needle (20) pipettes reagents from reagent containers (44) into reaction chambers. A container rack (90) keeps reagent containers securely in place.

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**METHODS AND APPARATUS FOR THE  
GENERATION OF CHEMICAL LIBRARIES**

**5    FIELD OF THE INVENTION**

The present invention relates generally to combinatorial synthesis, and more particularly to methods and apparatus for the generation of chemical libraries of known composition.

**10    BACKGROUND**

The relationship between structure and function of molecules is a fundamental issue in the study of biological and other chemistry-based systems. Structure-function relationships are important in understanding, for example, the function of enzymes, cellular communication, and cellular control and feedback mechanisms. Certain macromolecules are known to interact and bind to other molecules having a specific three-dimensional spatial and electronic distribution. Any macromolecule having such specificity can be considered a receptor, whether the macromolecule is an enzyme, a protein, a glycoprotein, an antibody, an oligonucleotide sequence of DNA, RNA or the like. The various molecules to which receptors bind are known as ligands.

Pharmaceutical drug discovery is one type of research that relies on the study of structure-function relationships. Much contemporary drug discovery involves discovering novel ligands with desirable patterns of specificity for biologically important receptors. Thus, the time necessary to bring new drugs to market could be greatly reduced through the use of methods and apparatus which allow rapid generation and screening of large numbers of ligands.

A common way to generate such ligands is to synthesize libraries of ligands on solid phase resins. Techniques for solid phase synthesis of peptides are described, for example, in Atherton and Sheppard; Solid

Phase Peptide Synthesis: A Practical Approach, IRL Press at Oxford University Press, Oxford, England, 1989.

Techniques for solid phase synthesis of oligonucleotides are described in, for example, Gait, Oligonucleotide

- 5 Synthesis: A Practical Approach, IRL Press at Oxford University Press, Oxford, England, 1984. Each of these references is incorporated herein by reference.

- Techniques for solution and solid phase multiple component combinatorial array syntheses strategies include U.S. Patent Application No. 08/092,862 filed Jan. 10 13, 1994, which is assigned to the assignee of the present invention, and which is incorporated herein by reference. Other synthetic strategies that may be employed are described in, for example, Bunin and Ellman, 15 "A General and Expedient Method for the Solid Phase Synthesis of 1,4-Benzodiazepine Derivatives," J. Amer. Chem. Soc. 114:10997-10998 (1992); Bunin et al., "The Combinatorial Synthesis and Chemical and Biological Evaluation of a 1,4-Benzodiazepine Library," Proc. Natl. Acad. Sci. 91:4708-4712 (1994); U.S. Patent No. 5,288,514 20 entitled "Solid Phase and Combinatorial Synthesis of Benzodiazepine Compounds on a Solid Support," issued Feb. 22, 1994; and PCT Publication WI 94/08051, April 14 (1994), each of which is incorporated herein by 25 reference.

- Since the introduction of solid phase synthesis methods for peptides, oligonucleotides and other polynucleotides, new methods employing solid phase strategies have been developed that are capable of 30 generating thousands, and in some cases even millions, of individual peptide or nucleic acid polymers using automated or manual techniques. These synthesis strategies, which generate families or libraries of compounds, are generally referred to as "combinatorial 35 chemistry" or "combinatorial synthesis" strategies.

Combinatorial chemistry strategies can be a powerful tool for rapidly finding novel ligands to receptors of interest. To date, three general strategies for generating combinatorial libraries have emerged:

- 5 "spatially-addressable," "split-bead," and "recombinant" strategies. These methods differ in one or more of the following aspects: reaction vessel design, polymer type and composition, control of physical variables such as time, temperature and atmosphere, isolation of products, 10 solid-phase or solution-phase methods of assay (i.e., chemical analysis), simple or complex mixtures, and methods for finding or determining the structure of the individual library members.

- Of these general strategies, several sub-strategies 15 have been developed. One spatially-addressable strategy that has emerged involves the generation of peptide libraries on immobilized pins that fit the dimensions of standard, 96 well microtiter plates. See PCT patent publication Nos. 91/17271 and 91/19818, each of which is 20 incorporated herein by reference. This method has been used to identify peptides which mimic discontinuous epitopes. Geysen et al., "Screening Chemically Synthesized Peptide Libraries for Biologically Relevant Molecules," Bioorg. Med. Chem. Lett. 3: 397-404 (1993), 25 and to generate benzodiazepine libraries, U.S. Patent No. 5,288,514 entitled "Solid Phase and Combinatorial Synthesis of Benzodiazepine Compounds on a Solid Support," issued Feb. 22, 1994 and Bunin et al., "The Combinatorial Synthesis and Chemical and Biological 30 Evaluation of a 1,4-Benzodiazepine Library," Proc. Natl. Acad. Sci. 91:4708-4712 (1994). The structures of the individual library members can be determined by analyzing the pin location (in the microtiter plate) in conjunction with the sequence of reaction steps (called a "synthesis 35 histogram") performed during the synthesis.

A second, related spatially-addressable strategy that has emerged involves solid-phase synthesis of polymers in individual reaction vessels, where the individual vessels are arranged into a single reaction unit. An illustrative example of such a reaction unit is a standard 96-well microtiter plate; the entire plate comprises the reaction unit and each well corresponds to a single reaction vessel. This approach is an extrapolation of traditional single-column solid-phase synthesis.

As is exemplified by the 96-well plate reaction unit, each reaction vessel is spatially defined by a two-dimensional matrix. Thus, the structures of individual library members can be determined by analyzing the sequence of reactions to which each well was subjected.

Another spatially-addressable strategy employs "teabags" (i.e., small, porous sacks) to hold synthesis resin. The reaction sequence to which each teabag is subject is recorded. This recorded reaction sequence determines the structure of the oligomer synthesized in each teabag. See for example, Lam *et al.*, "A New Type of Synthetic Peptide Library for Identifying Ligand-Binding Activity," *Nature* 354:82-84 (1991), Houghton *et al.*, "Generation and Use of Synthetic Peptide Combinatorial Libraries for Basic Research and Drug Discovery," *Nature* 354:84-86 (1991), and Jung *et al.*, "Multiple Peptide Synthesis Methods and Their Applications," *Angew. Chem. Int. Ed. Engl.* 31:367-383 (1992), each of which is incorporated herein by reference.

In another recent development, the techniques of photolithography, chemistry and biology have been combined to create large collections of oligomers and other compounds on the surface of a substrate. See U.S. Patent No. 5,143,854 and PCT patent publication Nos. 90/15070 and 92/10092, each of which is incorporated

herein by reference.

Recombinant methods for preparing collections of oligomers have also been developed. See PCT patent publication nos. 91/17271 and 91/19818, each of which is incorporated herein by reference. Using these methods, one can identify each oligomer in the library by determining the DNA coding sequences in a recombinant organism or phage. However, since the library members are generated in vivo (i.e., within the organism or phage), recombinant methods are limited to polymers whose synthesis can occur in the cell. Thus, these methods typically have been restricted to constructing peptide libraries.

A third general strategy that has emerged involves the use of "split-bead" combinatorial synthesis strategies. See Furka et al., "General Methods for Rapid Synthesis of Multicomponent Peptide Mixtures," Int. J. Pept. Protein Res. 37: 487-493, (1991) which is incorporated herein by reference. In this method, beads are apportioned into smaller groups. These smaller groups (called "aliquots") each contain a number of beads that is evenly divisible into the total number of beads. Each aliquot exposed to a monomer, and the beads are pooled together again. The beads are mixed, reapportioned into aliquots, and then exposed to a second monomer. The process is repeated until the desired library is generated.

A technique for synthesizing labelled combinatorial chemistry libraries is described in co-pending application serial number 08/383,776, entitled "Methods and Apparatus for Synthesizing Labeled Combinatorial Chemical Libraries," filed February 2, 1995, assigned to the assignee of the present invention, and incorporated herein by reference. In a preferred embodiment of that invention, each synthesized compound is associated with a unique identifier tag. The identifier tag relates a

signal to a detector upon excitation with electromagnetic radiation.

To aid in the generation of combinatorial chemical libraries, scientific instruments have been produced ---  
5 which automatically perform many or all of the steps required to generate such libraries. An example of an automated combinatorial chemical library synthesizer is the Model 396 MPS fully automated multiple peptide synthesizer, manufactured by Advanced ChemTech, Inc.  
10 ("ACT") of Louisville, KY.

The Model 396 MPS is capable of generating up to 96 different peptides or other small molecules in a single run. The syntheses occur simultaneously, with one amino acid being added to each growing polypeptide chain before  
15 addition of the next successive amino acid to any polypeptide chain. Thus, each growing polypeptide chain contains the same number of amino acid residues at the end of each synthesis cycle.

The syntheses occur in an ACT proprietary plastic reaction block having 96 reaction chambers. While the  
20 ACT reaction blocks work for their intended purpose, they possess several shortcomings.

First, ACT reaction blocks are machined from a single piece of plastic. Thus, they require extremely  
25 intricate machining, and are quite expensive to manufacture. Furthermore, since ACT reaction blocks are in the form of a single unit, should a portion of a block become damaged or contaminated in some way, the entire reaction block would have to be discarded; there is no  
30 way to replace individual portions of an ACT block.

An additional drawback of the plastic ACT reaction blocks is that they cannot be efficiently heated or cooled to aid in chemical reactions that may require such heating or cooling.

35 Certain processes and chemistries require that the chemical reagents (which may be reactants, solvents, or



reactants dissolved in solvents) be kept under an inert or anhydrous atmosphere to prevent reactive groups from reacting with molecular oxygen, water vapor, or other agents commonly found in air. Examples of atmosphere or moisture sensitive chemistries include peptide chemistry, nucleic acid chemistry, organometallic, heterocyclic, and chemistries commonly used to construct combinatorial chemistry libraries. Accordingly, such reagents must be stored and used under an anhydrous or inert atmosphere, such as one of argon, nitrogen, or other gases or mixtures of gases. Typically, containers of such reagents (and containers in which reactions using these reagents take place) are sealed from outside air by a gas impermeable septum. Reagents may be removed from or introduced into a septum sealed container via a non-coring pipetting needle that pierces the septum.

The composition of the septum depends on the chemistry involved, but common materials include thermoplastic rubber (TPR), natural rubber, teflon (typically used as a lining), and EPDM.

While the ACT reaction block can maintain an inert atmosphere when locked in place on the work station of the Model 396 MPS, there is no way to maintain an inert atmosphere once an ACT reaction block is removed from the work station. Thus, the reaction block must remain docked at the work station during the entire synthesis cycle. Since many reactants require several hours to react, this represents significant down time for the Model 396 MPS pipetting station, as it remains idle during the reaction period.

Additionally, creating an effective seal that maintains an inert atmosphere within the ACT reaction block is difficult due to the design of the block. To create such a seal, a top plate fitted with a rubber gasket is clamped onto the reaction chamber using six set screws. The screws are hand tightened to create the

seal. The top of the block is machined such that a raised rim or bead separates the 96 reaction chambers into four sections of 24 reaction chambers each. Thus, individual chambers within a group of 24 are not sealed with a raised bead but rather sealed with a flat junction between the septa and the flat top of the machined polymeric reaction block. This design provides an inferior seal and allows solvent from the reaction chambers to cross contaminate reaction chambers within each group of 24 by creeping along the underside of the septa material or alternatively, by creeping along the gas passages machined into the top of the reaction block. Proper adjustment of the screws to distribute pressure evenly across each of the four sections (to create an effective seal) requires careful manipulation and cannot always be accomplished successfully.

A poorly formed seal can also create a problem with reagent cross-contamination. If the gasket does not seal evenly around each reaction chamber, reagents may seep from one reaction vessel into another.

While the ACT reaction block includes 96 reaction chambers, the compounds generated in the ACT reaction block cannot be directly transferred into a standard 96-well microtiter plate because the distance between the outlets of the reaction chambers is too great. For each reaction chamber to have the volume needed to perform reactions, the 96-reaction chamber ACT reaction block must necessarily be too large to mate with a standard 96-well microtiter plate. When reactions are complete, the user must transfer the contents of the reaction chambers into an array of 96 flat bottom glass vials supported in a plastic frame. The user must then manually pipette fluid from the glass vials into a microtiter plate for further analysis. This arrangement presents several disadvantages. First, the glass vials must be cleaned between uses, which increases the chance for

contamination. Furthermore, the labor intensive nature of the transfer increases a chance for error. Finally, ~~this process cannot easily be automated.~~

The reagent delivery system of the Model 396 MPS  
5 also suffers limitations. While the septum-sealed reagent containers from which the reactants are drawn can be sealed under an inert or anhydrous atmosphere, the volume of reagent removed is not replaced with an equivalent or greater volume of inert gas. As reagents  
10 are withdrawn from the reagent containers, a partial vacuum is generated within the containers. If the pressure difference between the inside of the container and the external atmosphere is great enough, outside air may seep into the container through needle holes  
15 previously made in the septum.

The Model 396 MPS also employs a capacitance detector that can determine the fluid surface level in a reagent bottle. During operation of the Model 396 MPS, fluid is removed from reagent bottles by inserting a  
20 pipetting needle just below the fluid surface level such that reagent directly at the reagent-atmosphere interface is withdrawn. While this operation permits only the very tip of a pipetting needle to be contaminated, this operation may also result in the withdrawal of reagent  
25 that has been exposed to outside air.

Finally, the MPS detector described above can only operate if polar reagents are used. Thus, the Model 396 MPS may not be compatible with chemistries that utilize non-polar reagents.

30 Accordingly, there remains a need in the art for a relatively inexpensive, easy to manufacture reaction block having replaceable parts. There also remains a need in the art for a reaction block that can be efficiently heated and cooled, that can be moved from  
35 place to place while maintaining an inert atmosphere, and that can mate directly with a standard 96 well microtiter

plate. An additional need that remains in the art is for a reaction block that easily and effectively seals each reaction chamber, and that reduces cross-contamination of reagents between reaction chambers. There also remains a  
5 need in the art for a reaction block that can be manipulated robotically.

There also remains a need in the art for a pipetting work station that can be operated to withdraw reagent from the bottom of a reagent bottle, away from the  
10 reagent-atmosphere interface, and that can be used with non-polar reagents.

There remains a further need in the art for a holder for septa sealed reagent bottles which prevents the movement of these bottles caused by friction between a  
15 pipetting needle and the septa seals.

#### SUMMARY

The preferred embodiments meet these needs by providing a reaction block which uses replaceable  
20 reaction chambers supported in the block. Each reaction block is fitted with four sets of 12 reaction chambers, and has fittings that facilitate robotic manipulation.

The reaction chambers are preferably fitted with a frit. An s-shaped trap tube snaps into a fitting on the  
25 bottom of each reaction chamber. The trap tube runs into a drain tube.

The reaction block is fitted with gas (preferably N<sub>2</sub>) lines and a septum seal such that gas pressurization empties the reaction chambers into the drain tubes. The  
30 drain tubes are arranged so as to mate directly with a standard 96 well microtiter plate for the collection of material.

The reaction blocks preferably have high thermal stability and are preferably fitted with gas or liquid  
35 lines for heating or cooling.

A docking station provides for secure registration

of the reaction blocks, and provides for introduction of gases and liquids into the reaction blocks. An inert atmosphere in the reaction block is maintained by a top and (optional) bottom seal. A synthesis support may be introduced as a slurry or powder through the fastened septa with a suitable septa-piercing needle assembly. Alternatively, a synthesis support may be introduced into each reaction chamber as a slurry or powder with the top removed and the top then fastened after addition. A needle pipettes reagents from an array of reagent containers into the reaction chambers, and maintains the inert atmosphere. A locking reagent container rack keeps the containers securely in place.

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**BRIEF DESCRIPTION OF THE DRAWINGS**

**Figure 1** is a plan view of a pipetting work station according to a preferred embodiment.

**Figure 2** is a side and top view of a reaction chamber according to a preferred embodiment.

**Figure 3** is a side cross-sectional view of the reaction chamber shown in **Figure 2** further including a trap tube and drain tube.

**Figure 4** is an isometric view of a reaction block and its associated hardware according to a preferred embodiment.

**Figure 5** is a top view of the reaction block shown in **Figure 4**.

**Figure 6** is a side cross-sectional view of the reaction block shown in **Figure 4**.

**Figure 7** is a side cross-sectional view of the reaction block shown in **Figure 4** including a removable bottom seal.

**Figure 8** is a side cross-sectional view of the reaction block shown in **Figure 4** including a microtiter plate.

**Figure 9** is a bottom view of the reaction block shown in **Figure 4**.

**Figure 10** is a bottom plan view of the reaction block shown in **Figure 4**.

**Figure 11** is a top view of a locking reagent container rack according to a preferred embodiment.

**Figure 12** is a side view of the locking reagent container rack shown in **Figure 11**.

**Figure 13** is a top view of a docking station according to a preferred embodiment.

**Figure 14** is a cross sectional view of a connector in the docking station shown in **Figure 13**, inserted into a port in the reaction block shown in **Figure 4**.

**Figure 15** is a cross sectional view of a connector in the docking station shown in **Figure 13**, inserted

into a port having a open valve in the reaction block shown in **Figure 4**.

**Figure 16** is a cross sectional view of a connector in the docking station shown in **Figure 13**, inserted  
5 into a port having a closed valve in the reaction block shown in **Figure 4**.

#### DETAILED DESCRIPTION

The structure and function of the preferred  
10 embodiments can best be understood by reference to the drawings. The reader will note that the same reference numerals appear in multiple figures. Where this is the case, the numerals refer to the same or corresponding structure in those figures.

#### 15 GENERAL OPERATION

**Figure 1** is a plan view showing a portion of an automated pipetting work station **250** as may be used in a preferred embodiment. Automated pipetting work station **250** may be a TECAN 5032 automated pipetting work station  
20 (manufactured by TECAN AG, Feldbachstrasse 80, CH-8634 Hombrechtikon, Switzerland) with one or more pipetting arms **252**. Pipetting arm **252** attaches to needle assembly **20**, which is preferably a coaxial needle assembly of the type disclosed in concurrently filed application serial  
25 no. 08/423,142 (attorney docket no. 8140-010) entitled "Pipetting Needle for Fluid Transfer Under Inert Atmosphere Operations," assigned to the assignee of the present invention and incorporated herein by reference. Coaxial needle assembly **20** includes a needle **22**, a gas  
30 inlet port **30**, and may also include an electrical connection **31**. Work station **250** may also include pipetting needle rinse stations **70**, preferably of the type disclosed in concurrently filed application serial  
no. 08/423,141 (attorney docket no. 8140-011) entitled  
35 "Pipetting Needle Rinse Station" assigned to the assignee of the present invention, and incorporated herein by

reference.

A locking reagent container rack 90 holds several containers 44 of reagents sealed from the outside air with septum seals 46. Rack 90 is preferably placed on the left side of work station deck 254. On the right side of work station deck 254 is a docking station 300 for receiving two reaction blocks 140. Each reaction block 140 contains an array of 48 reaction chambers 110 (see, e.g., Figure 4). A standard 96 well microtiter plate 302 may be mounted below reaction block 140 when product is to be removed from reaction chambers 110.

#### REACTION CHAMBER

Referring now to Figure 2, a reaction chamber 110 according to a preferred embodiment is shown. Reaction chamber 110 is preferably made of an injection molded or extruded polymer such as polypropylene, although polyethylene, teflon, glass, or any other inert material able to withstand the temperature, pressure, and chemical environment to which reaction chamber 110 is exposed could also be used. Reaction chamber 110 preferably also has an internal volume of approximately 2 ml.

Reaction chamber 110 includes a generally cylindrical body portion 112 of a first diameter, and a generally cylindrical lower body portion 114 that is coaxial with and of a smaller diameter than body portion 112. Body portion 112 is connected to generally cylindrical lower body portion 114 by frustum, or tapered section 116. Body portion 112 has a top opening 118 which preferably has a rounded top surface 120 to facilitate sealing against a septum. Body 112 has a gas input port 122 located below top surface 120. Gas input port 122 is positioned such that it will be located above the bottom of waste basin 160 of reaction block 140 (See Figures 4 and 5). As will be discussed below, this positioning will prevent possible cross-contamination of chemicals by blocking a direct flow path



from port to port or from port to septum.

Reaction chamber 110 preferably also includes a  
keying protrusion 128. Keying protrusion 128 prevents  
reaction chamber 110 from being inserted into reaction  
5 block 140 unless it is in a predetermined orientation.  
This feature will be discussed further below.

Lower portion 114 of reaction chamber 110 can  
receive a frit 124, which preferably supports a quantity  
of a synthesis support such as solid phase resin (not  
10 shown). Frit 124 is preferably a 70 micron polyethylene  
frit, although other types of frits such as sintered  
glass, sintered metals, and sintered ceramics may be used  
depending on the type of chemistry to be performed.

Frit 124 is preferably press fit into lower portion  
15 114. Lower portion 114 has a smaller diameter than that  
of body 112 to allow insertion of frit 124 into lower  
portion 114 without damaging the inside surfaces of body  
112. Lower portion 114 preferably also includes an  
annular bead 126 to retain frit 124 when it is pressed  
20 into place.

Reaction chamber 110 also includes a funnel shaped  
portion 130 immediately below lower portion 114, and  
below frit 124. Funnel shaped portion 130 allows full  
area exposure to the underside of frit 124, to enhance  
25 the draining of liquids from reaction chamber 110.

Immediately below funnel portion 130 is a generally  
cylindrical drain tube 132. Drain tube 132 is coaxial  
with and of smaller diameter than lower body portion  
114. Drain tube 132 includes an annular sealing bead  
30 134 to create a seal against the outside of S-shaped  
trap tube 136 (see **Figure 3**). This seal may be  
strengthened by welding (or otherwise fixing) S-shaped  
trap tube 136 to annular sealing bead 134.

The purpose of trap tube 136 is to prevent the loss  
35 of liquids from reaction chamber 110 (when reaction  
chamber 110 is not pressurized) by bringing the level of

an outlet for liquid above the normal liquid level of reaction chamber 110. Trap tube 136 connects to a drain tube 138. When reaction chamber 110 is pressurized, liquid flows through trap tube 136 and out drain tube 138. As will be discussed below, drain tube 138 will be positioned so as to deposit liquid into a well of a standard 96-well microtiter plate.

Trap tube 136 and drain tube 138 are preferably made of an injection molded or extruded polymer such as polypropylene, although polyethylene, teflon, or any other inert material able to withstand the temperature, pressure, and chemical environment to which trap tube 136 and drain tube 138 are exposed could also be used.

#### REACTION BLOCK

Referring now to Figure 4, an isometric view of a reaction block 140 (and its associated hardware) according to a preferred embodiment is shown. Reaction block 140 is preferably machined out of 6061 aluminum (which is easily machinable and has good corrosion resistance) and then anodized for additional corrosion protection. Reaction block 140 could also be hard coat anodized followed by teflon impregnation. Additionally reaction block 140 could be machined or molded from any suitable metal, engineering plastics, filled plastics, crystalline plastics, ceramics, machinable ceramics, or any other material that can withstand the temperature, pressure, and chemical environment to which reaction block 140 will be exposed. If non-metallic materials are used, product reaction could be enhanced by the application of microwaves. If materials transparent to ultraviolet (UV) light are used, product could be cleaved from the synthesis support using UV light, and without the application of an acid or base.

Each end of reaction block 140 is preferably fitted with two pins 178 to facilitate handling by a robotic gripper (not shown). Each side of reaction block 140 is

preferably fitted with one pin 180 to facilitate securing reaction block 140 onto docking station 300. Robotic manipulation of reaction block 140 makes automation of the entire synthesis process possible. For example, reagents could be introduced into reaction chambers 110 when reaction block 140 is locked onto docking station 300 of pipetting work station 250. Reaction block 140 could then be moved to a separate docking station 300, vortexing shaker table, heating or cooling chamber, or any other location or device (not shown) useful in synthesis or the collection of material.

In a preferred embodiment, two types of reaction blocks capable of mating directly with a 96 well microtiter plate are contemplated: the 48 reaction chamber 110 (and drain tube 138) positions of a first type of (or "A") block are offset from the 48 reaction chamber and drain tube positions of a second type of (or "B") block such that a type "A" and a type "B" block can fill every position in a standard 96 well microtiter plate. The ability to deposit material directly into a 96-well microtiter plate eliminates the possible contamination and human error problems discussed above with respect to the ACT reaction block.

Reaction block 140 may be color coded for ease of identification, may have identification numbers 320 machined into or printed on the sides, and may also have a bar code 322 printed on the side for identification by machine.

Referring now to Figure 5, top portion 142 of reaction block 140 is shown. Top portion 142 preferably has an array of circular openings 144 arranged in a staggered grid. In a preferred embodiment, reaction block 140 has 48 circular openings 144. Openings 144 also preferably include a keying notch 145 (see Figure 4) which cooperates with keying protrusion 128 on reaction chamber 110 and requires reaction chamber 110

to be in a predetermined orientation when inserted into opening 144.

The 48 openings 144 are divided into four chambers 146A through 146D of twelve openings 144 each. The chambers 146A-D are separated from each other by a plurality of raised beads 148, which are preferably machined into top portion 142.

Top portion 142 also includes four gas inlet ports 150A through 150D. Gas flows from gas inlet ports 150A-D, into gas inlet chambers 152A through 152D, respectively (which are defined by raised sealing beads 148). Gas then flows out through chamber exit ports 154A through 154D, respectively, and into chambers 146A through 146D, respectively. Gas flow to each chamber 146 can be individually controlled. For example, chambers 146A and 146C can be pressurized, without pressurizing chambers 146B and 146D.

When reaction chambers 110 are inserted and locked into place in openings 144, the top portions 120 of reaction chambers 110 are in approximately the same plane as the tops of raised sealing beads 148.

Top surfaces 120 of reaction chambers 110 and raised sealing beads 148 are sealed by a sheet of septum material 153 (See Figure 4). Septum 153 is preferably manufactured from 1/10" thermoplastic rubber (TPR) sheet.

Septum 153 is retained by a septum retainer plate 155, which is preferably fastened with six captive screw-type fasteners 156 which attach to openings 157. Fasteners 156 pass through openings 159 in septum 153, and screw into machined fastener openings 158.

Reaction block 140 may be sealed from underneath with a bottom seal 220. An o-ring or quad ring 221 (see Figure 7) may be used to ensure a gas-tight seal. Bottom seal 220 may include a one-way valve 222 to allow pressure regulation. Bottom seal 220 is preferably fitted to reaction block 140 with screw-type fasteners

224. As can be seen in **Figure 4**, fasteners **224** pass through openings **226** in plate **155**, through openings **228** in septum **153**, through openings **228** in reaction block **140**, and into openings **232** in bottom seal **220**. Bottom seal **220** permits a desired atmosphere or pressure to be maintained within reaction block **140**, allowing reaction block **140** to be moved from location to location (such as to a separate shaker table, not shown) without loss of such atmosphere or pressure. This can be especially useful in chemistries requiring long periods of time for reactions to take place. In these situations, such reactions can take place away from the pipetting work station, allowing the pipetting work station to be used for other purposes.

In a preferred embodiment, septum retainer plate **155** is machined from 6061 aluminum and anodized. However, retainer plate **155** could also be machined or molded from engineering plastics, ceramics, or any other material that can withstand the temperature, pressure, and chemical environment to which retainer plate **155** will be exposed.

Plate **155** is also preferably machined with **48** openings **162** positionally matched with openings **144** of reaction block **140** (and thus with openings **118** of reaction chambers **110**) to accurately control the compression of the septum **153** between the tops **120** of reaction chambers **110**, and plate **155**.

Chambers **146 A through D** include recessed waste basins **160 A through D**, respectively, which are machined into top portion **142** below the level of chamber exit ports **154A - D**. This prevents a back flow of fluids from waste basins **160A - D** into chamber exit ports **154A - D**.

As discussed above, reaction chambers **110** and openings **144** are preferably "keyed" with keying protrusions **128** and keying notches **145**, respectively.

This prevents reaction chambers 110 from being inserted fully into openings 144 unless the reaction chambers are in a predetermined proper orientation. In a preferred embodiment, reaction chambers 110 are oriented such that gas inlet ports 122 face away from chamber exit ports 154A - D. This prevents back flow of liquids from reaction chambers 110 into the chamber exit ports 154A - D. In addition, gas inlet ports 122 of reaction chambers 110 are oriented such that a back flow of liquid from one reaction chamber 110 is prevented from spilling directly into the gas inlet port 122 of an adjoining reaction chamber 110.

Referring now to **Figure 6**, a side cross-sectional view of reaction block 140 is shown. Reaction chambers 110 are held in place by machined annular steps 170 (which define openings 171), and machined annular beads 172. S-shaped trap tube 136 and drain tube 138 are held in place by a friction fit against walls 174 and openings 176 (See **Figures 9 and 10**).

Steps 177 are machined into the bottom of reaction block 140 to allow reaction block 140 to mate directly with a standard 96-well microtiter plate 302 (see, e.g., **Figures 1 and 8**). Steps 177 also allow mating and sealing with bottom seal 220 (see **Figures 4 and 7**).

Referring now to **Figures 9 and 10**, plan and bottom views of reaction block 140 are shown. The underside of reaction block 140 includes a generally planar surface 190 which includes a plurality of openings 171 and 176, discussed above. Openings 176 include a relatively larger portion 192, which accommodates drain tube 138, and a relatively smaller portion 194, which accommodates s-shaped trap tube 136.

The underside of reaction block 140 also includes four gas ports 196A through 196D located on bottom surface 198. Ports 196A-D connect to gas inlet ports 150A-D (See **Figure 5**), respectively, through machined

tunnels (not shown) in reaction block 140.

Also included on bottom surface 198 is a gas inlet port 200 which connects to a gas outlet port 201 via a machined tunnel (not shown). This allows pressure on the underside of reaction block 140 to be independently controlled when it is sealed by bottom seal 220 (see Figures 4 and 7).

Bottom surface 198 also includes two gas or liquid ports 202A and 202B. The interior of reaction block 140 is preferably machined to include passages (not shown) in which heating or cooling gas or liquid can flow if desired. Gas or liquid can enter port 202A and exit through port 202B, or vice versa. If reaction block 140 is made of material having high thermal stability or thermal mass such as 6061 aluminum, this arrangement allows reaction block 140 to be quickly and efficiently heated or cooled for chemistries that require such heating or cooling.

Ports 196A-D, 200 and 202 may also serve as guide pin holes to position reaction block 140 properly on docking station 300 (see Figures 1 and 13).

Finally, a bar magnet 204 may be mounted flush with surface 198. Bar magnet 204 serves to activate magnetic reed switch 314 mounted in docking station 300 (see Figure 13). As will be discussed below, one or more reed switches prevent the operation of work station 250 unless one or more reaction blocks 140 are properly in place.

#### **DOCKING STATION**

Referring now to Figures 1, 13 and 14, a docking station 300 according to a preferred embodiment is shown. Docking station 300 preferably includes two stations, 306A and 306B, for receiving reaction blocks 140 of Type "A" and Type "B", respectively, as discussed above. As is known to those skilled in the art, docking station 300 may also be fitted with the proper motor,

gears, and other elements (not shown) necessary for docking station 300 to act as a vortexing shaker, and preferably as a vortexing shaker having a fixed displacement and variable speed.

5        Docking station 300 also preferably includes three locking linkages 304 for locking onto pins 180 on reaction blocks 140. Each station 306 includes gas outlet connectors 308A through 308D which connect to ports 196A through 196D, respectively in reaction block  
10    140 (see **Figure 9**). Each station 306 also includes two coolant or heating fluid (i.e., gas or liquid) connectors 310A and 310B. **Figure 1** shown fluid lines 320A and 320B attached to connectors 310A and 310B, respectively. Although not shown in **Figures 1** and **13**,  
15    independently controllable fluid lines attach to each connector shown in docking station 300. Connectors 310A and 310B connect to ports 202A and 202B, respectively in reaction block 140 (See **Figure 9**). A gas outlet connector 312 which connects to gas inlet port 200 of  
20    reaction block 140 is also included in each station 306.

Finally, stations 306A and 306B each include a magnetic reed switch 314A and 314B, respectively, which senses the presence of magnet 204 on reaction block 140.

Station 306A, and more specifically the placement of port 312, is arranged such that only an A-type reaction  
25    block 140 can be fully inserted and locked into position. Similarly, station 306B, and more specifically the placement of port 312, is arranged such that only a B-type reaction block 140 can be fully  
30    inserted and locked into position.

Referring now to **Figure 14**, a cross sectional view of a connector 308A inserted into port 196A of reaction block 140 is shown. Although only the interface between connector 308A and 196A will be discussed, it will be  
35    understood that similar interfaces are preferably included in other connections between reaction block 140



and docking station 300. In a preferred embodiment, connector 308A is inserted into port 196A. In this fashion, connector 308A acts as a guide pin to ensure proper alignment of reaction block 140 with station 306A. A gas-tight seal between connector 308A and port 196A is preferably provided by quad ring 330. A quad ring is preferred over a standard o-ring because a quad ring has less tendency to adhere to surfaces when connector 308A is removed from port 196A.

Referring now to Figures 15 and 16 an alternative embodiment of port 196A is shown. In operations where inert or other atmosphere must be maintained, a normally closed valve, such as schraeder valve 360 may be placed in port 196A. Schraeder valve 360 may be replaced with a bi-directional elastomeric valve. In operation, connector 308A is inserted into port 196A and engages pin 362 of schraeder valve 360. Connector 308A also forms a seal against quad ring 330. Gas flows out of opening 364 and through schraeder valve 360.

When connector 308A is removed from port 196A, pin 362 of schraeder valve 360 moves downward, creating a gas-tight seal. Again, this allows reaction block 140 to be moved from place to place while maintaining a desired atmosphere.

Pipetting work station 250 (see Figure 1) is preferably constructed such that operation of the work station cannot take place unless magnetic reed switches 314A and 314B detect the presence of one or both reaction blocks 140. That is, pipetting work station 250 will not operate unless reaction blocks 140 are properly mounted in stations 306A and 306B.

#### **LOCKING REAGENT CONTAINER RACK**

As was discussed above, pipetting reagents under inert atmosphere is often essential during the synthesis of combinatorial libraries. However, when pipetting reagents from a relatively lightweight septum-sealed

container, the friction between the pipetting needle and the septum may be enough to lift the container from its resting position. This is obviously not desirable, as movement of and damage to the container and other equipment may result. Referring now to **Figures 1, 11, and 12**, a preferred embodiment of a locking reagent container rack 90 for pipetting under inert atmosphere is shown. Rack 90 includes a bottom plate or grating 92, preferably coated with a layer of rubber or soft material 94. Rack 90 may include one or more horizontal support plates 96, and includes a plurality of vertical support members 98. Horizontal plates 96 preferably include 48 circular openings 97 into which the containers 44 can be inserted. Rack 90 is preferably arranged such that septum-sealed containers 44 can be arranged in a 4 x 8 array.

A top plate 100 rests on top of vertical support members 98 and snugly against containers 44. Top plate 100 preferably includes 48 circular openings 102 into which the tops of containers 44 can be inserted. Each opening 102 is preferably encircled with a rubber ring 104 to protect containers 44.

Plate 100 also includes one or more fasteners (such as quarter turn wing-nut type fasteners) 105 which fasten plate 100 to vertical support members 98, thus keeping containers 44 in place. Locking rack 90 may itself be fastened to a work surface (such as work station deck 254), although the weight of rack 90 and containers 44 would probably be sufficient to prevent any motion caused by friction between a pipetting needle and a septum.

#### **EXAMPLE OF OPERATION**

The many features of the preferred embodiments described above facilitate the relatively quick and efficient generation of chemical libraries. In the following discussion, a synthesis operation involving a

type "A" reaction block 140 will be discussed. However, it will be understood that the following discussion will apply equally for a type "B" block as well.

In a typical operation, a synthesis support such as solid phase resin is deposited onto each frit 124 in reaction chambers 110. Reaction block 140 is then assembled as shown in Figure 4. Bottom seal 220 may be mounted if reaction block 140 must be moved from place to place while maintaining a desired atmosphere or pressure.

Reaction block 140 may then be manually or robotically inserted into station 306A of docking station 300 on work station 250 (see Figures 1 and 13). At this point, microtiter plate 302 is not located in station 306A. Locking linkages 304 then grip pins 180, locking reaction block 140 into place. A type "B" reaction block may be simultaneously mounted in station 306B.

Pipetting work station 250 then operates under computer control to deliver the chosen combination of reagents into reaction chambers 110. Specifically, pipetting needle 22 (as controlled by pipetting arm 252) is used to transfer reagents from septum 46 sealed containers 44 into septum 253 sealed reaction chambers 110. The interior and exterior of pipetting needle 22 may be cleaned as necessary in rinse stations 70. At any time that reaction block 140 is mounted in station 306A, reaction block 140 may be heated or cooled, pressurized with inert gas, or vortexed as described above.

For reactions that take a considerable amount of time, reaction block 140 may be manually or robotically moved to another docking station 300, or to some other location while the reactions are taking place. After the synthesis of the desired products has been completed, the products may be cleaved from the synthesis supports using

the appropriate reagents. These reagents may be applied at work station 250, or they may be applied robotically at some other location. If bottom seal 220 had been mounted, it is then removed, and reaction block 140 is mounted onto a microtiter plate 302 in station 306A. Reaction chambers 110 are then pressurized, forcing the product out drain tubes 138 and into alternate wells of microtiter plate 302. Microtiter plate 302 is then moved to station 306B. A type "B" reaction block 140 is mounted on microtiter plate 302, and product is then deposited into the alternate empty wells of microtiter plate 302 as discussed above. Again, this process allows product to be deposited directly into the wells of a standard microtiter plate, without requiring an intermediate step.

The present invention has been described in terms of a preferred embodiment. The invention, however, is not limited to the embodiment depicted and described. Rather, the scope of the invention is defined by the appended claims.

## WHAT IS CLAIMED IS:

1. A reaction chamber comprising:
  - 5 a generally cylindrical upper body having a top opening, an inner surface, an outer surface, and an inside diameter;
  - a generally cylindrical lower body coaxial with the upper body;
  - 10 the lower body having an inner surface, an outer surface, and an inside diameter smaller than the inside diameter of the upper body;
  - a tapered section connecting the upper body to the lower body; and
  - 15 ~~a drain tube having an inside surface and~~ connected to the lower body.
2. A reaction chamber as in claim 1 wherein the drain tube is connected to the lower body with a funnel-shaped section.
- 20 3. A reaction chamber as in claim 1 wherein the top opening of the upper body has a rounded edge to facilitate sealing against a septum.
- 25 4. A reaction chamber as in claim 1 wherein the upper body includes a port connecting the inner surface of the upper body to the outer surface of the upper body.
- 30 5. A reaction chamber as in claim 4 wherein the port is a gas inlet port.
- 35 6. A reaction chamber as in claim 1 wherein the drain tube is connected to an s-shaped trap tube having an inlet opening and an outlet opening.

7. A reaction chamber as in claim 6 wherein the drain tube has an annular bead on its inside surface to facilitate sealing with the inlet opening of the s-shaped trap tube.
- 5 8. A reaction chamber as in claim 7 wherein the s-shaped trap tube is welded to the drain tube.
- 10 9. A reaction chamber as in claim 6 wherein the s-shaped trap tube extends upwardly along the length of the upper body.
- 15 10. A reaction chamber as in claim 9 wherein the outlet port of the s-shaped trap tube points downward.
- 20 11. A reaction chamber as in claim 10 wherein the outlet port of the s-shaped trap tube is connected to a second cylindrical drain tube.
- 25 12. A reaction chamber as in claim 11 wherein the second cylindrical drain tube is of a larger diameter than the trap tube.
- 30 13. A reaction chamber as in claim 12 wherein the s-shaped trap tube and the second cylindrical drain tube are made from extruded polypropylene or teflon.
- 35 14. A reaction chamber as in claim 1 wherein the upper body, the lower body, the tapered section, the funnel shaped section, and the drain tube are made from a single piece of injection-molded polymer.
15. A reaction chamber as in claim 14 wherein the polymer is polypropylene or teflon.

16. A reaction chamber as in claim 1 wherein the upper body has a volume of approximately 2 ml.
- 
17. A reaction chamber as in claim 1 wherein the upper body further includes a keying protrusion on its outer surface.
18. A reaction chamber as in claim 1 wherein the lower body further includes a frit having a diameter smaller than the inside diameter of the lower body portion.
- 
19. A reaction chamber as in claim 18 wherein the ~~frit is press-fit into the lower body portion.~~
- 
20. A reaction chamber as in claim 19 wherein the frit is made of polyethylene.
21. A reaction chamber comprising:  
a generally cylindrical upper body having a top opening, an inner surface, an outer surface, and an inside diameter;  
a generally cylindrical lower body coaxial with the upper body;  
the lower body having an inner surface, an outer surface, and an inside diameter smaller than the inside diameter of the upper body;  
means for connecting the upper body to the lower body;  
a drain tube having an inside surface; and means for connecting the drain tube to the lower body.
22. A reaction chamber as in claim 21 wherein the means for connecting the drain tube to the lower

body is a funnel-shaped section.

23. A reaction chamber as in claim 21 wherein the  
means for connecting the upper body to the lower  
body is a tapered section.
24. A reaction chamber as in claim 21 wherein the  
top opening of the upper body has a rounded edge to  
facilitate sealing against a septum.
25. A reaction chamber as in claim 21 wherein the  
upper body includes a means for connecting the inner  
surface of the upper body to the outer surface of  
the upper body.
26. A reaction chamber as in claim 25 wherein the  
means for connecting the inner surface of the upper  
body to the outer surface of the upper body port is  
a gas inlet port.
27. A reaction chamber as in claim 21 wherein the  
drain tube is connected to an s-shaped trap tube  
having an inlet opening and an outlet opening.
28. A reaction chamber as in claim 27 wherein the  
drain tube has an annular bead on its inside surface  
to facilitate sealing with the inlet opening of the  
s-shaped trap tube.
29. A reaction chamber as in claim 28 wherein the  
s-shaped trap tube is welded to the drain tube.
30. A reaction chamber as in claim 27 wherein the  
s-shaped trap tube extends upwardly along the length  
of the upper body.



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31. A reaction chamber as in claim 30 wherein the outlet port of the s-shaped trap tube points downward.
- 5 32. A reaction chamber as in claim 31 wherein the outlet port of the s-shaped trap tube is connected to a second cylindrical drain tube.
- 10 33. A reaction chamber as in claim 32 wherein the second cylindrical drain tube is of a larger diameter than the trap tube.
- 15 34. A reaction chamber as in claim 33 wherein the s-shaped trap tube and the second cylindrical drain tube are made from extruded polypropylene or teflon.
- 20 35. A reaction chamber as in claim 21 wherein the upper body, the lower body, the tapered section, the funnel shaped section, and the drain tube are made from a single piece of injection-molded polymer.
36. A reaction chamber as in claim 35 wherein the polymer is polypropylene or teflon.
- 25 37. A reaction chamber as in claim 21 wherein the upper body has a volume of approximately 2 ml.
38. A reaction chamber as in claim 21 wherein the upper body further includes a keying protrusion on its outer surface.
- 30 39. A reaction chamber as in claim 21 wherein the lower body further includes means for retaining a frit.
- 35 40. A reaction chamber as in claim 39 wherein the frit is press fit into the lower body portion.

41. A reaction chamber as in claim 40 wherein the frit is made of polyethylene.

- 5 42. A reaction chamber comprising:  
a generally cylindrical upper body having  
a top opening with a rounded edge to facilitate  
sealing against a septum, an inner surface, an  
outer surface, a gas port connecting the inner  
10 surface with the outer surface, and an inside  
diameter;  
a generally cylindrical lower body  
coaxial with the upper body;  
the lower body having an inner surface  
15 with an annular bead for retaining a frit, an  
outer surface, and an inside diameter smaller  
than the inside diameter of the upper body;  
a tapered section connecting the upper  
body to the lower body;  
20 a drain tube having an inside surface;  
and a funnel shaped section connecting  
the drain tube to the lower body.
- 25 43. A reaction block comprising:  
a first side wall and a second side wall,  
the first and second side walls having a first  
width and a first height;  
a first end wall and a second end wall,  
the first and second end walls having a second  
30 width narrower than the first width, and the  
first height;  
a top surface bordered by the first and  
second side walls and the first and second end  
walls, the top surface having:  
35 a plurality of generally circular  
openings for removably receiving a plurality of

reaction chambers, the reaction chambers each having a circular opening and a top surface;

a gas port;

and a plurality of raised beads defining the edges of a plurality of chambers;

the raised beads having a top surface which is in approximately the same plane as the top surface of the reaction chambers;

a flexible septum which seals against the top surfaces of the reaction chambers and the raised beads;

and a retainer plate which covers the septum.

44. A reaction block as in claim 43 wherein the retainer plate has a plurality of openings which spatially correspond to the openings of the reaction chambers.
45. A reaction block as in claim 44 wherein the retainer plate is held securely against the septum and the reaction block with a plurality of fasteners.
46. A reaction block as in claim 43 wherein the top surface includes a first, a second, a third, and a fourth, separately controllable gas inlet ports, and wherein the raised beads define a first, a second, a third, and a fourth pressurization chambers.
47. A reaction block as in claim 46 wherein each pressurization chamber includes an equal number of the reaction chambers.
48. A reaction block as in claim 46 wherein each pressurization chamber includes 12 reaction chambers.

49. A reaction block as in claim 46 wherein the first, second, third, and fourth separately controllable gas inlet ports are in communication with the first, second, third, and fourth pressurization chambers, respectively.
50. A reaction block as in claim 47 wherein each of the pressurization chambers includes a waste basin which is recessed below the level of the top surfaces of the reaction chambers.
51. A reaction block as in claim 50 wherein each of the four pressurization chambers has 12 of the reaction chambers.
52. A reaction block as in claim 43 wherein each of the generally circular openings on the top surface have a keying notch, and wherein each of the reaction chambers have a keying protrusion.
53. A reaction block as in claim 43 wherein the block is made of aluminum.
54. A reaction block as in claim 53 wherein the block is made of 6061 aluminum.
55. A reaction block as in claim 53 wherein the aluminum has been anodized.
56. A reaction block as in claim 43 wherein the block is color coded.
57. A reaction block as in claim 43 wherein the block has an identification number.

58. A reaction block as in claim 43 wherein the septum is made from thermoplastic rubber.
59. A reaction block as in claim 58 wherein the  
5 septum is die cut with six registration holes and four access holes.
60. A reaction block as in claim 43 wherein the  
10 first and second side walls are fitted with one pin each to facilitate securing the block to a work station.
61. A reaction block as in claim 43 wherein the  
15 first and second end walls are fitted with two pins each to facilitate handling by a robotic gripper.
62. A reaction block as in claim 43 further  
including a bottom surface defined by the bottom  
edges of the first and second side walls and the  
20 first and second end walls, the bottom surface being parallel to the top surface.
63. A reaction block as in claim 43 further  
including a first, second, third and fourth gas  
25 inlet ports in communication with the first, second, third, and fourth gas outlet ports of the top surface, respectively.
64. A reaction block as in claim 63 wherein the  
30 first, second, third and fourth gas inlet ports are in the bottom surface.
65. A reaction block as in claim 43 further  
35 including an input port for receiving heating or cooling fluid.

66. A reaction block as in claim 65 wherein the input port for receiving heating or cooling fluid is in the bottom surface.
- 5 67. A reaction block as in claim 65 further including an output port for draining heating or cooling fluid.
- 10 68. A reaction block as in claim 67 wherein the output port for draining heating or cooling fluid is on the bottom surface.
- 15 69. A reaction block as in claim 68 further including a plurality of passages connecting the input port for receiving heating or cooling fluid to the output port for draining heating or cooling fluid.
- 20 70. A reaction block as in claim 62 wherein the bottom surface includes a magnet.
71. A reaction block as in claim 70 wherein the magnet is a bar magnet.
- 25 72. A reaction block as in claim 62 wherein each of the reaction chambers includes a drain tube.
- 30 73. A reaction block as in claim 72 further including a cavity extending upwardly from the bottom surface to a middle surface, the middle surface parallel to the top surface and the bottom surface.
- 35 74. A reaction block as in claim 73 wherein the middle surface has openings for receiving the drain tubes.

75. A reaction block as in claim 74 wherein the cavity includes a machined step to facilitate mating with a microtiter plate.
- 5 76. A reaction block as in claim 75 wherein the cavity includes a second machined step to facilitate mating with a bottom seal.
- 10 77. A reaction block as in claim 76 wherein the bottom seal includes an o-ring which seals against the second machined step.
- 15 ~~78. A reaction block as in claim 76 wherein the bottom seal includes a one-way pressure valve.~~
- 20 79. A reaction block as in claim 63 wherein the first, second, third and fourth gas inlet ports include a valve.
80. A reaction block as in claim 80 wherein the valve is a schraeder valve.
81. A reaction block as in claim 80 wherein the valve is a bi-directional elastomeric valve.
- 25 82. A reaction block comprising:  
a first side wall and a second side wall,  
the first and second side walls having a first  
width and a first height;  
a first end wall and a second end wall,  
the first and second end walls having a second  
width narrower than the first width, and the  
first height;  
a top surface bordered by the first and  
second side walls and the first and second end
- 30  
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walls, the top surface having:

a plurality of generally circular openings for removably receiving a plurality of reaction chambers, the reaction chambers each having a circular opening and a top surface;

a gas port; and

a plurality of raised beads defining the edges of a plurality of chambers;

the raised beads having a top surface which is in approximately the same plane as the top surface of the reaction chambers;

means for sealing against the top surfaces of the reaction chambers and the raised beads;

and a retainer plate which covers the means for sealing.

83. A reaction block as in claim 83 wherein the retainer plate has a plurality of openings which spatially correspond to the openings of the reaction chambers.

84. A reaction block as in claim 84 further including means for fastening the retainer plate to the means for sealing the reaction block.

85. A reaction block as in claim 83 wherein the top surface includes a first, a second, a third, and a fourth, separately controllable gas inlet ports, and wherein the raised beads define a first, a second, a third, and a fourth pressurization chambers.

86. A reaction block as in claim 86 wherein each of the four pressurization chambers includes an equal number of the reaction chambers.



87. A reaction block as in claim 87 wherein the first, second, third, and fourth separately controllable gas inlet ports are in communication with the first, second, third, and fourth pressurization chambers, respectively.
88. A reaction block as in claim 87 wherein each of the four pressurization chambers includes a waste basin which is recessed below the level of the top surfaces of the reaction chambers.
89. A reaction block as in claim 87 wherein each of the four chambers has 12 of the reaction chambers.
90. A reaction block as in claim 83 wherein each of the generally circular openings on the top surface have a keying notch, and wherein each of the reaction chambers have a keying protrusion.
91. A reaction block as in claim 83 wherein the block is made of aluminum.
92. A reaction block as in claim 92 wherein the block is made of 6061 aluminum.
93. A reaction block as in claim 92 wherein the aluminum has been anodized.
94. A reaction block as in claim 83 wherein the block is color coded.
95. A reaction block as in claim 83 wherein the block has an identification number.
96. A reaction block as in claim 83 wherein the block has a bar code.

97. A reaction block as in claim 83 wherein the means for sealing is a septum
- 5 98. A reaction block as in claim 98 wherein the septum is made from thermoplastic rubber.
99. A reaction block as in claim 99 wherein the septum is die cut with six registration holes and four access holes.
- 10 100. ~~A reaction block as in claim 83 wherein the first and second side walls are fitted with one pin each to facilitate securing the block to a work station.~~
- 15 101. A reaction block as in claim 83 wherein the first and second end walls are fitted with two pins each to facilitate handling by a robotic gripper.
- 20 102. A reaction block as in claim 83 further including a bottom surface defined by the bottom edges of the first and second side walls and the first and second end walls, the bottom surface being parallel to the top surface.
- 25 103. A reaction block as in claim 83 further including a first, second, third and fourth gas inlet ports in communication with the first, second, third, and fourth gas outlet ports of the top surface, respectively.
- 30 104. A reaction block as in claim 104 wherein the first, second, third and fourth gas inlet ports are in the bottom surface.
- 35 105. A reaction block as in claim 83 further

including an input port for receiving heating or cooling fluid.

106. A reaction block as in claim 106 wherein the  
5 input port for receiving heating or cooling fluid is in the bottom surface.

107. A reaction block as in claim 106 further  
10 including an output port for draining heating or cooling fluid.

108. A reaction block as in claim 108 wherein the  
output port for draining heating or cooling fluid is on the bottom surface.

15 109. A reaction block as in claim 109 further including a plurality of passages connecting the input port for receiving heating or cooling fluid to the output port for draining heating or cooling  
20 fluid.

110. A reaction block as in claim 103 wherein the bottom surface includes a magnet.

25 111. A reaction block as in claim 111 wherein the magnet is a bar magnet.

112. A reaction block as in claim 103 wherein each of the reaction chambers includes a drain tube.

30 113. A reaction block as in claim 113 further including a cavity extending upwardly from the bottom surface to a middle surface, the middle surface parallel to the top surface and the bottom  
35 surface.

114. A reaction block as in claim 114 wherein the middle surface has openings for receiving the drain tubes.
- 5 115. A reaction block as in claim 115 wherein the cavity includes a machined step to facilitate mating with a microtiter plate.
- 10 116. A reaction block as in claim 116 wherein the cavity includes a second machined step to facilitate mating with a bottom seal.
- 15 117. A reaction block as in claim 117 wherein the bottom seal includes an o-ring which seals against the second machined step.
118. A reaction block as in claim 117 wherein the bottom seal includes a one-way pressure valve.
- 20 119. A reaction block as in claim 104 wherein the first, second, third and fourth gas inlet ports include a quad ring seal.
- 25 120. A reaction block as in claim 98 wherein the septum is made from an elastomeric material.
121. A reaction block as in claim 98 wherein the septum is made from an elastomeric material coated with teflon.
- 30 122. A container rack for use in a pipetting work station, comprising:  
a bottom surface for supporting a  
plurality of septum sealed reagent containers;  
35 a horizontal plate having a plurality of  
circular holes for receiving the containers;

a locking top plate having a plurality of circular holes aligned with the circular holes in the horizontal plate, and at least one fastener;

5 a plurality of vertical support members which support the locking top plate and the horizontal plate;

and a means for fastening the top plate to the vertical support members.

10

123. A rack as in claim 123 wherein the circular openings of the top plate are lined with a soft material.

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124. A rack as in claim 124 wherein the soft material is rubber.

125. A docking station for receiving a reaction block having a plurality of gas inlet ports, comprising:

20

a platform including a cavity for removably receiving a reaction block;

a plurality of gas outlet connectors coupled to the gas inlet ports of the reaction block;

25

a reed switch which detects the presence of the reaction block; and

a linkage which locks the reaction block into the cavity.

30

126. A docking station as in claim 126 wherein the reaction block includes a bar magnet, and wherein the reed switch detects the presence of the bar magnet.

35

127. A docking station as in claim 126 further

including a heating or cooling fluid outlet connector coupled to a heating or cooling fluid inlet port on the reaction block.

5 128. A docking station as in claim 126 wherein the platform includes a first cavity of a first type for receiving a first reaction block of a first type, and a second cavity of a second type for receiving a second reaction block of a second type.

10 129. ~~A docking station as in claim 127 wherein the gas outlet connectors fit into the gas inlet ports of the reaction blocks.~~

15 130. A docking station as in claim 128 wherein the gas outlet connectors are sealed against the gas inlet ports with quad rings.

20 131. A docking station as in claim 126 wherein said platform acts as a vortexing shaker.

25 132. A docking station for receiving a reaction block having a plurality of gas inlet ports, comprising:  
a platform including means for removably receiving a reaction block;  
means for connecting a supply of gas to the gas inlet ports of the reaction block;  
means for detecting the presence of the reaction block;  
30 and means for locking the reaction block into the means for removably receiving the reaction block.

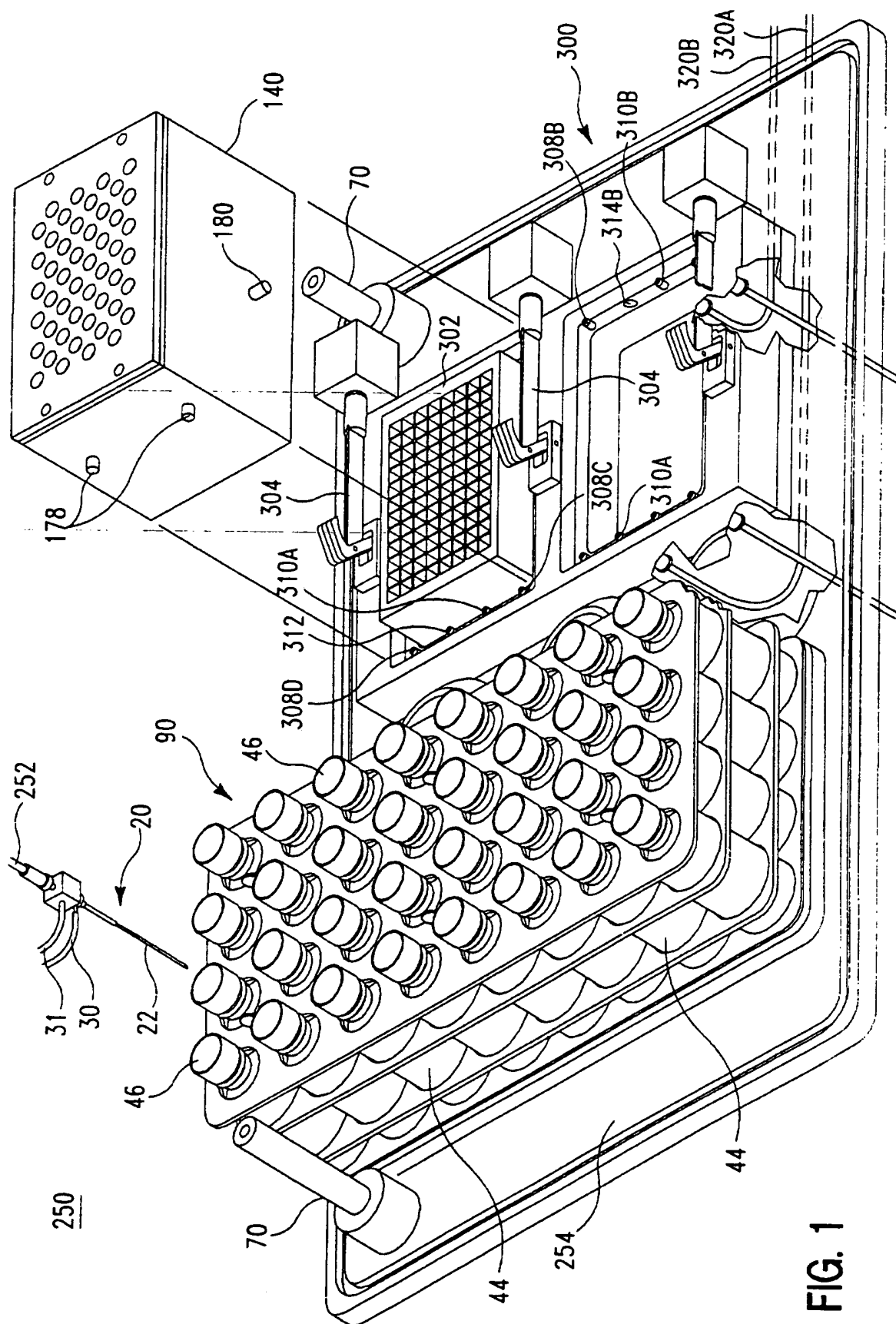
35 133. A docking station as in claim 133 wherein the reaction block includes a bar magnet, and wherein the means for detecting detects the presence of the

bar magnet.

134.       A docking station as in claim 133 further  
including means for connecting a source of heating  
5       or cooling fluid to a heating or cooling fluid inlet  
port on the reaction block.

135.       A docking station as in claim 133 wherein the  
platform includes a first means for receiving a  
10       first reaction block of a first type, and a second  
means for receiving a second reaction block of a  
second type.

136.       A docking station as in claim 133 wherein said  
15       platform acts as a vortexing shaker.





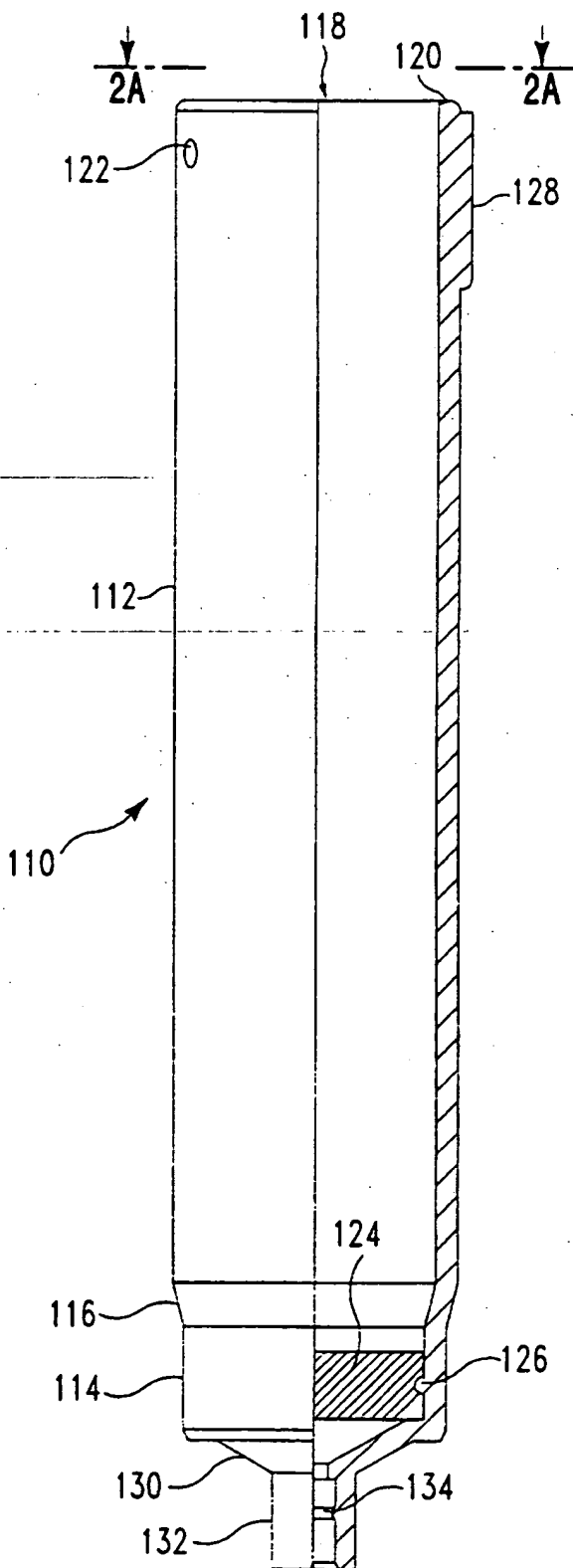
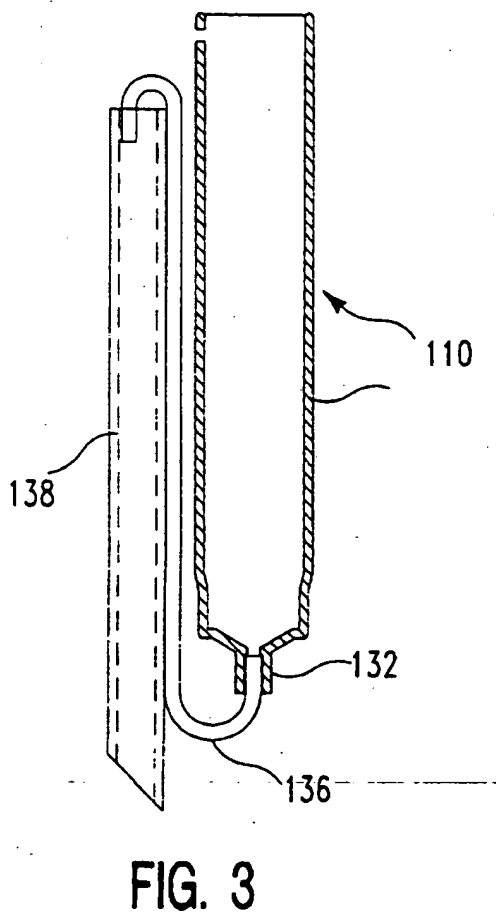
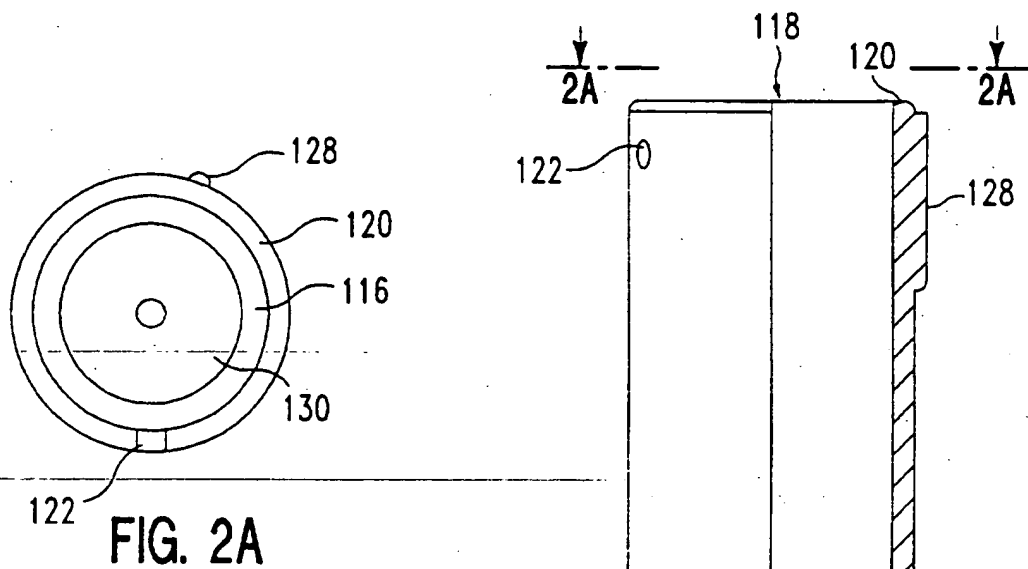


FIG. 2

FIG. 3

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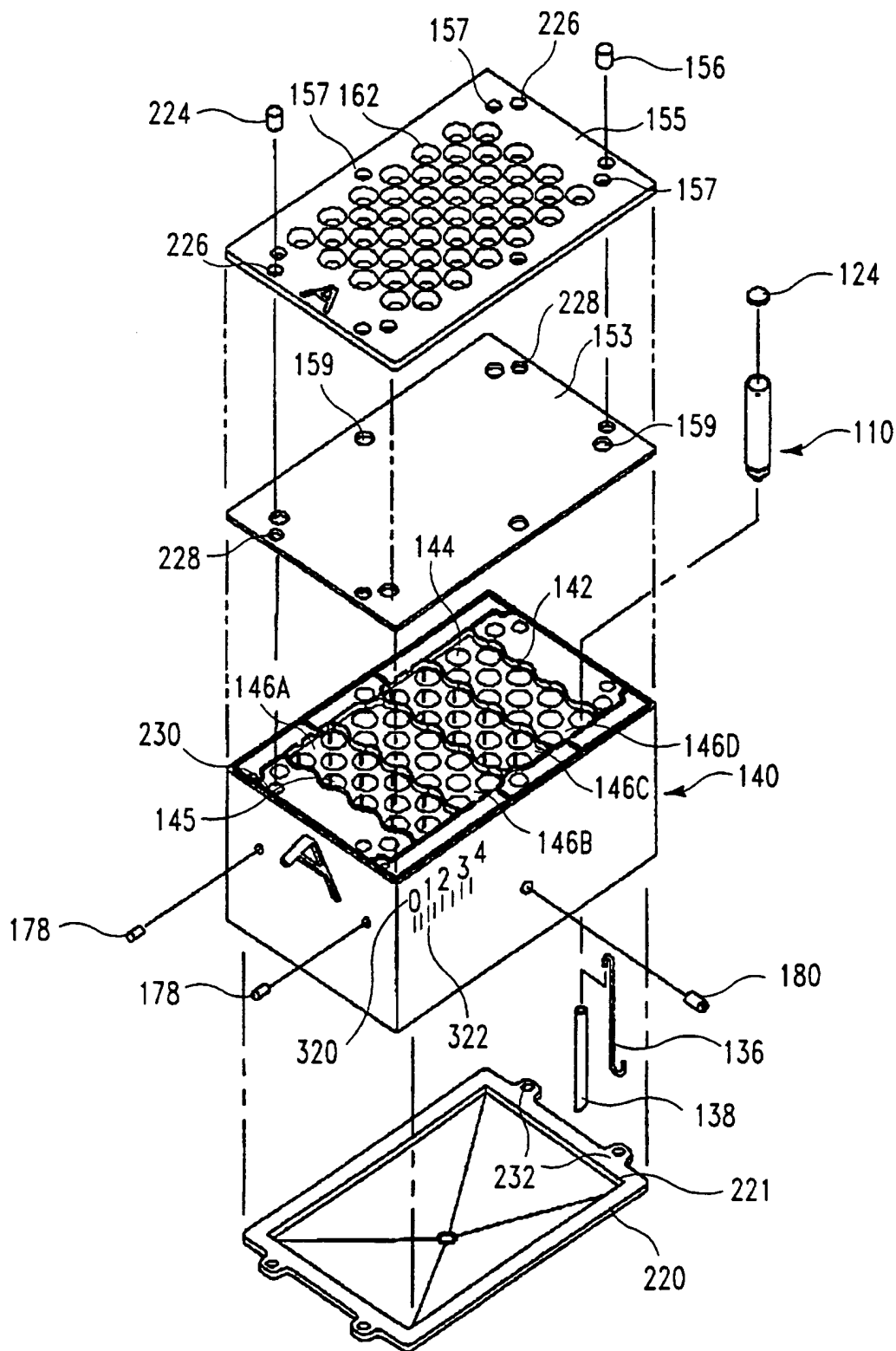
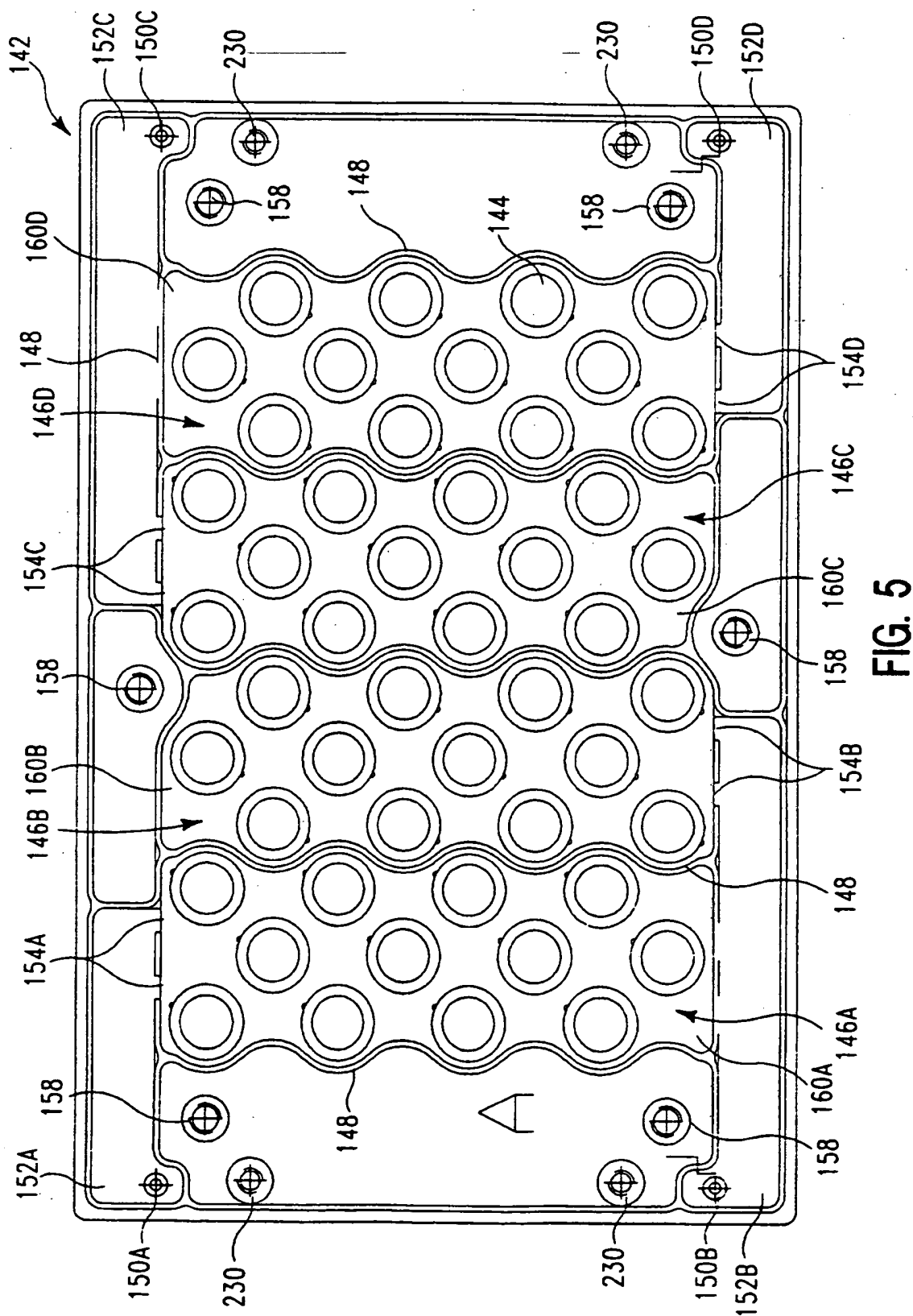


FIG. 4

SUBSTITUTE SHEET (RULE 28)



**SUBSTITUTE SHEET (RULE 26)**

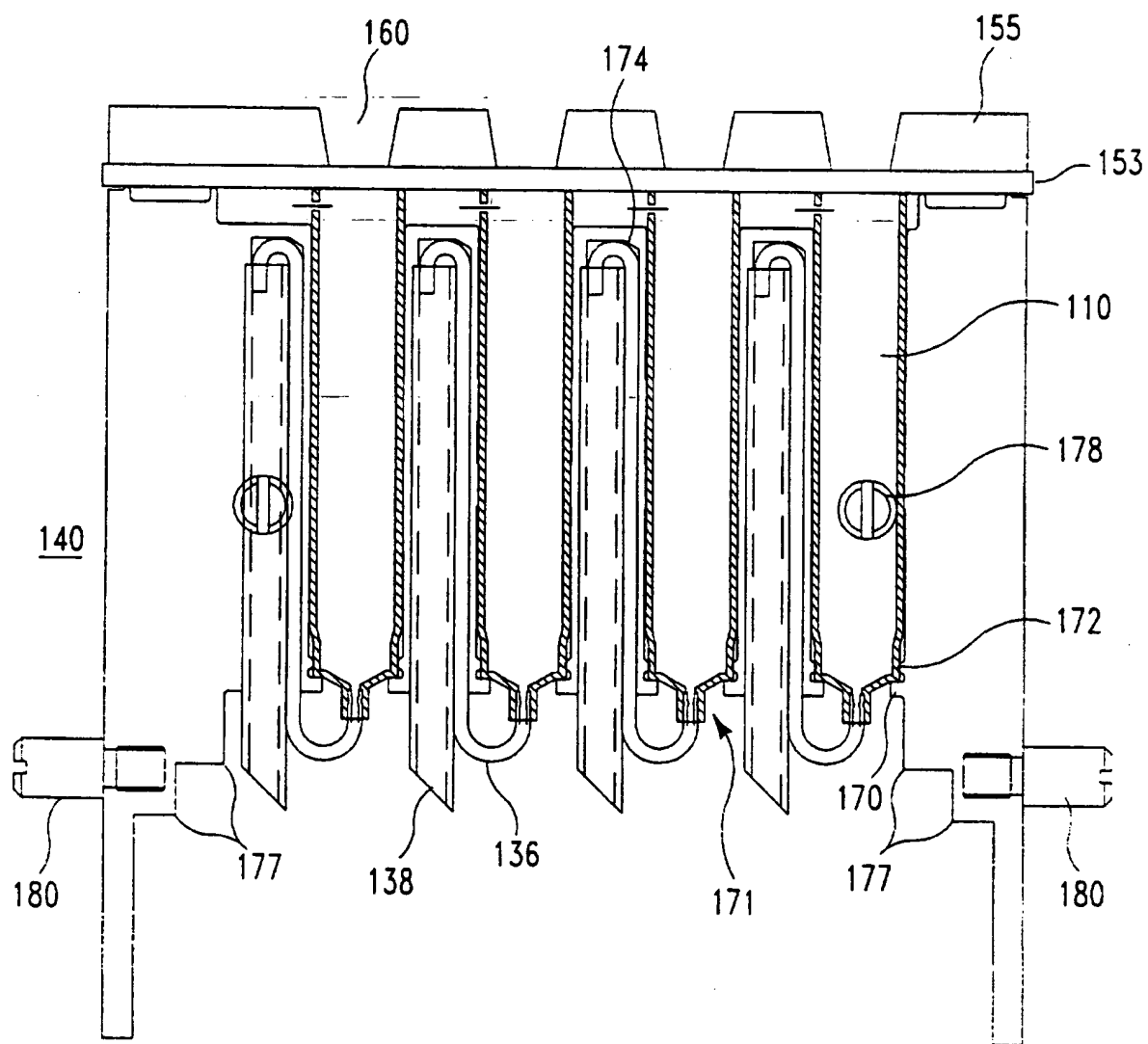


FIG. 6

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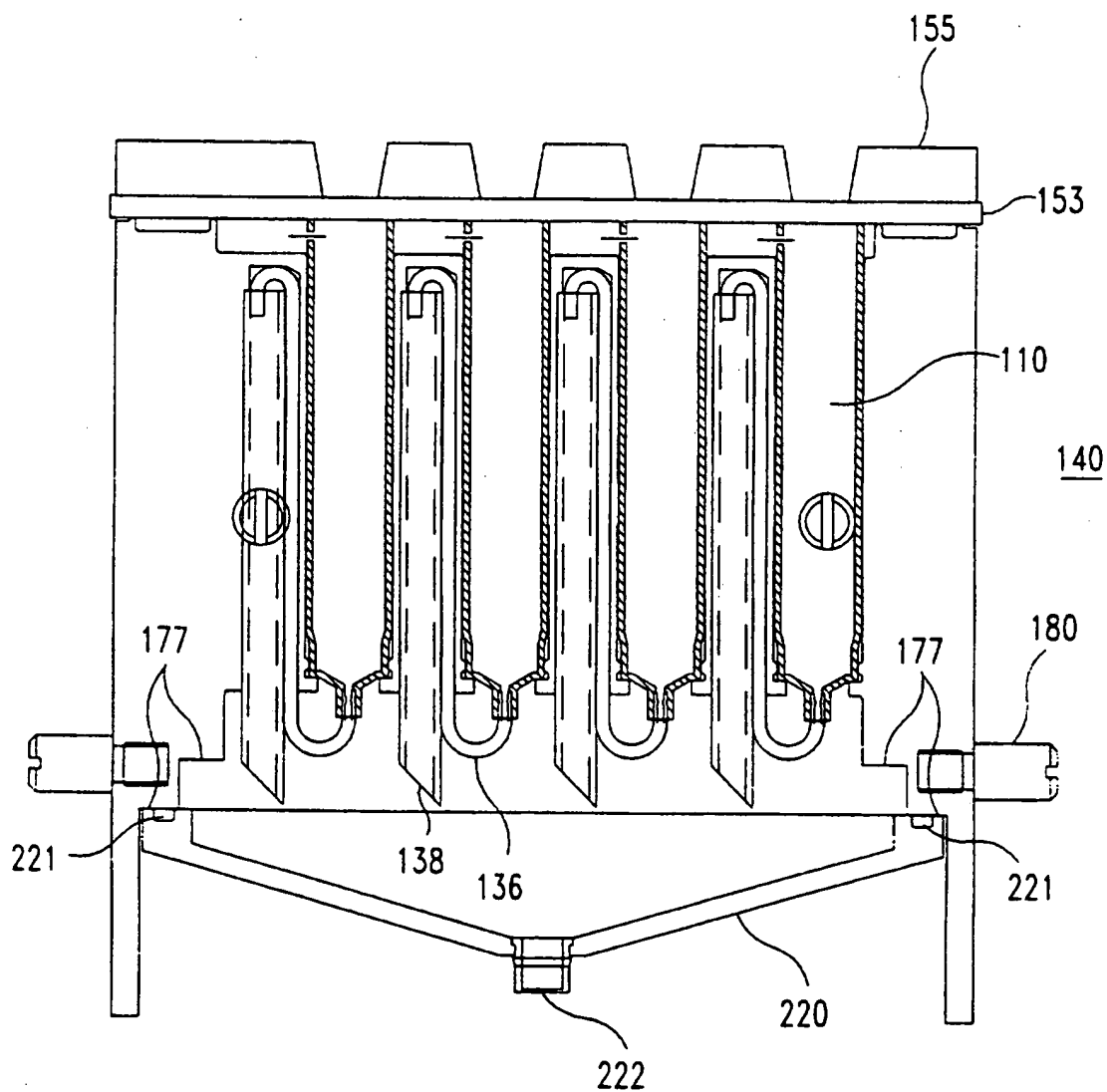


FIG. 7

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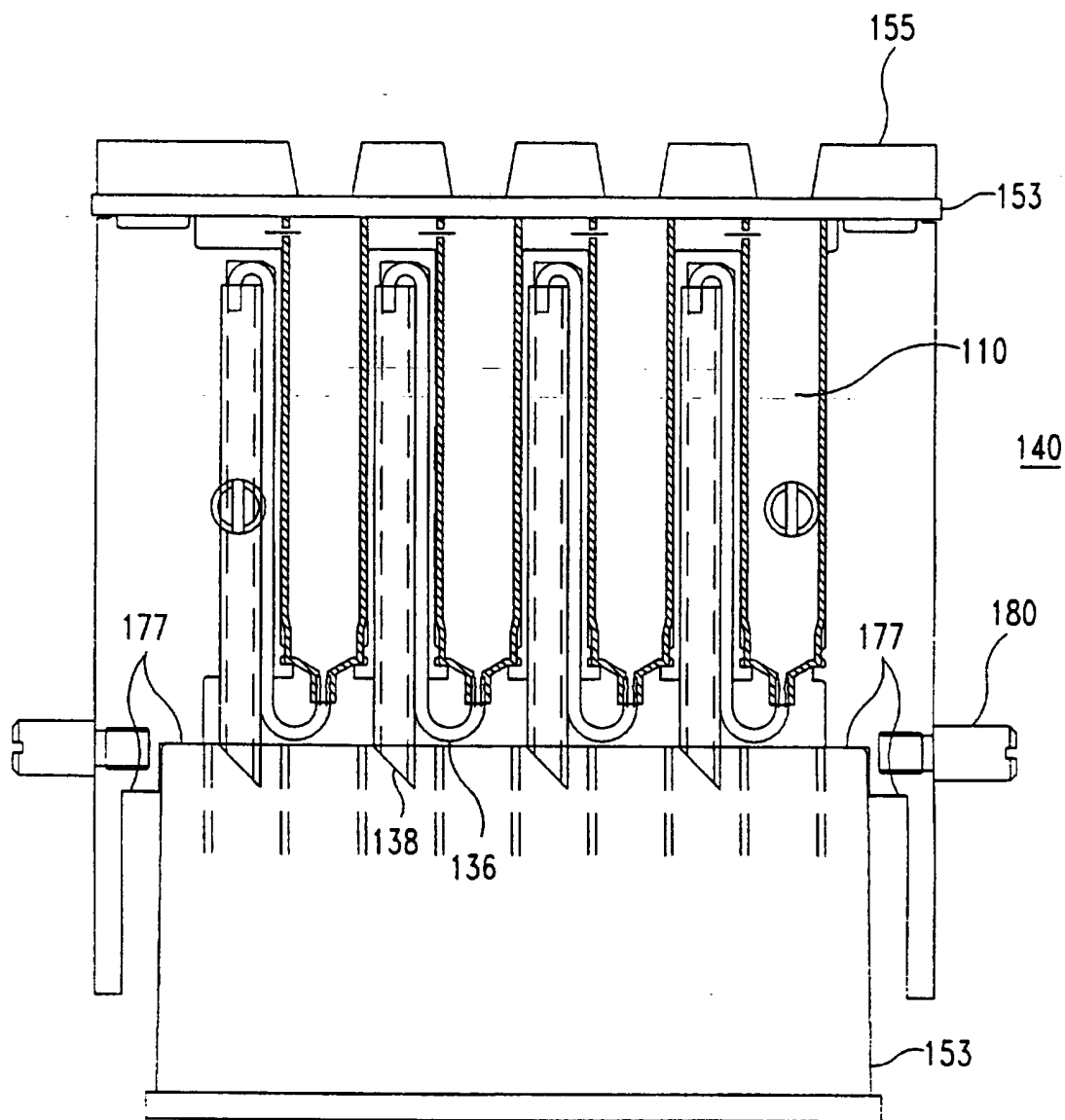


FIG. 8

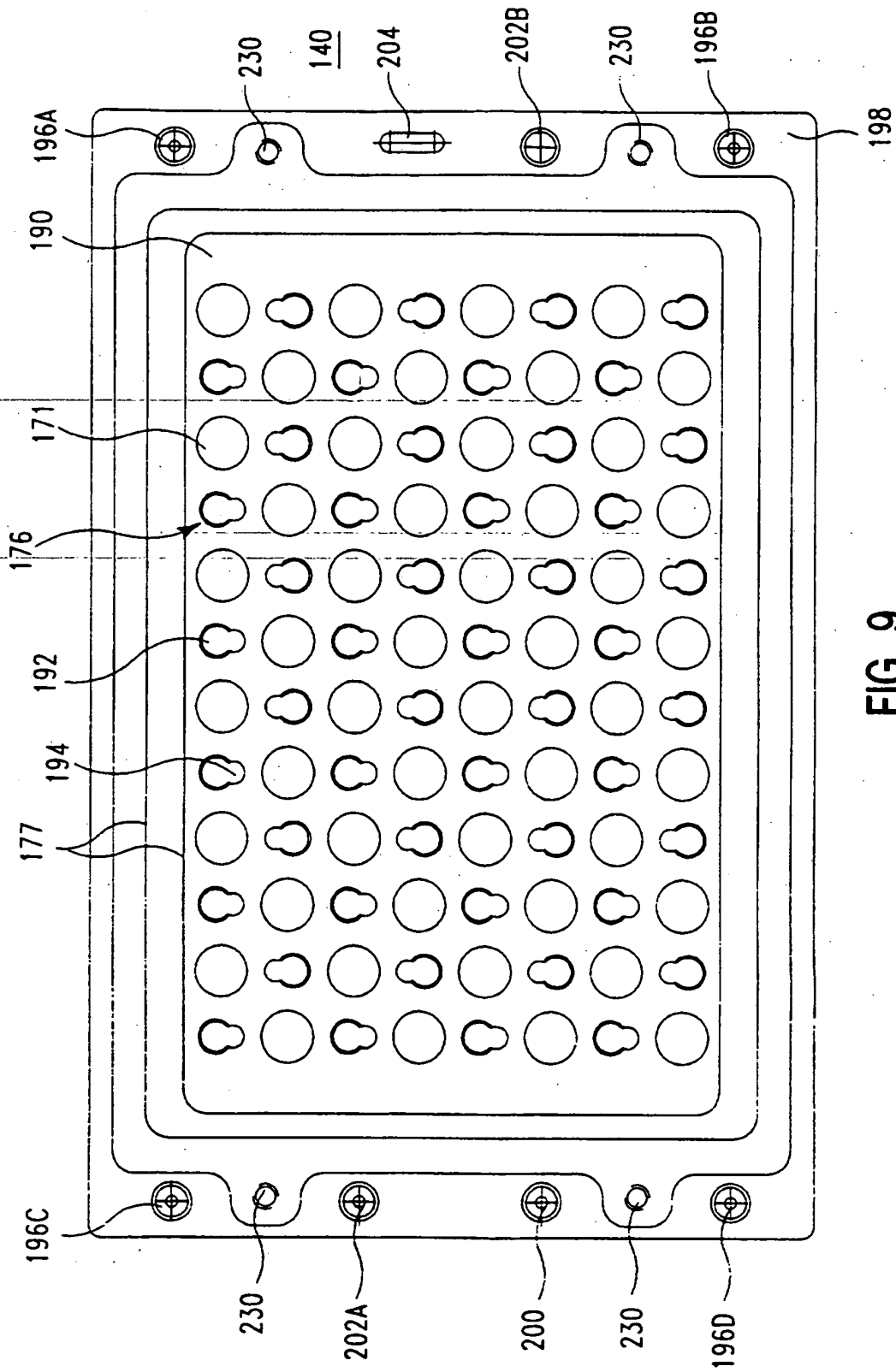


FIG. 9

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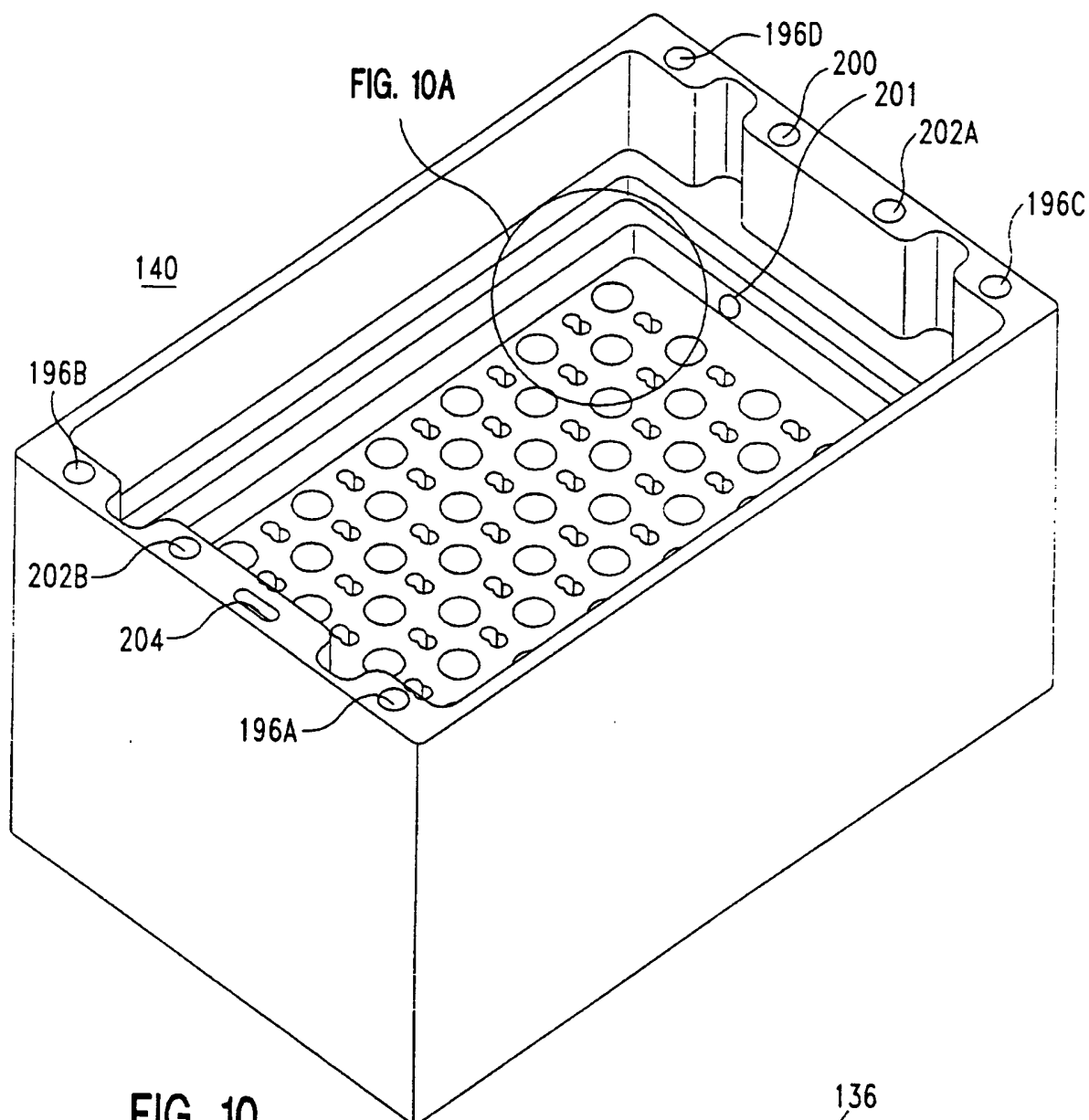
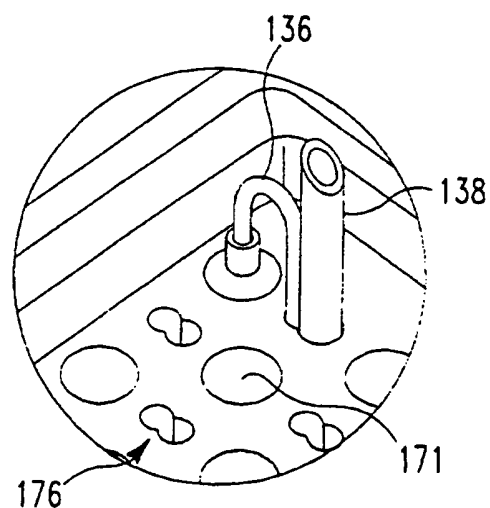


FIG. 10





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FIG. 11

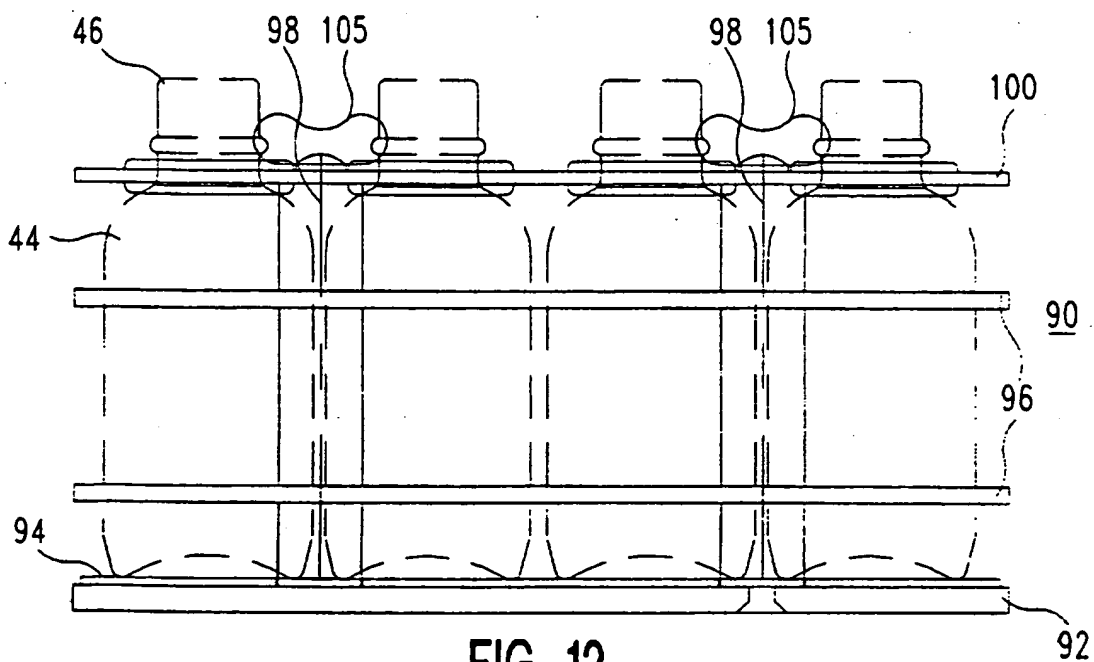
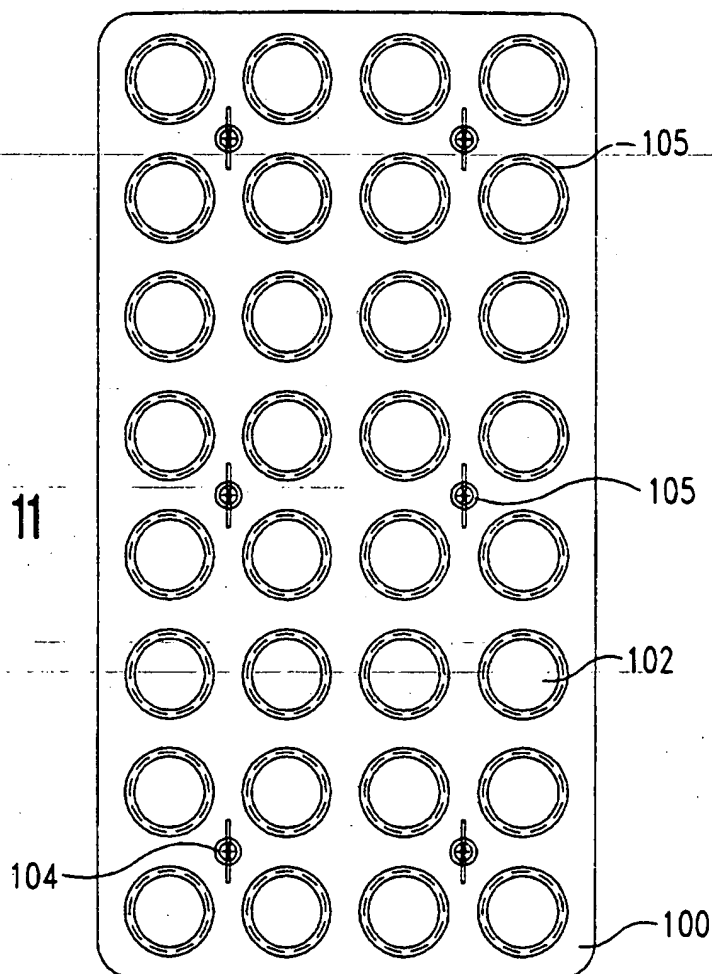


FIG. 12

SUBSTITUTE SHEET (RULE 26)

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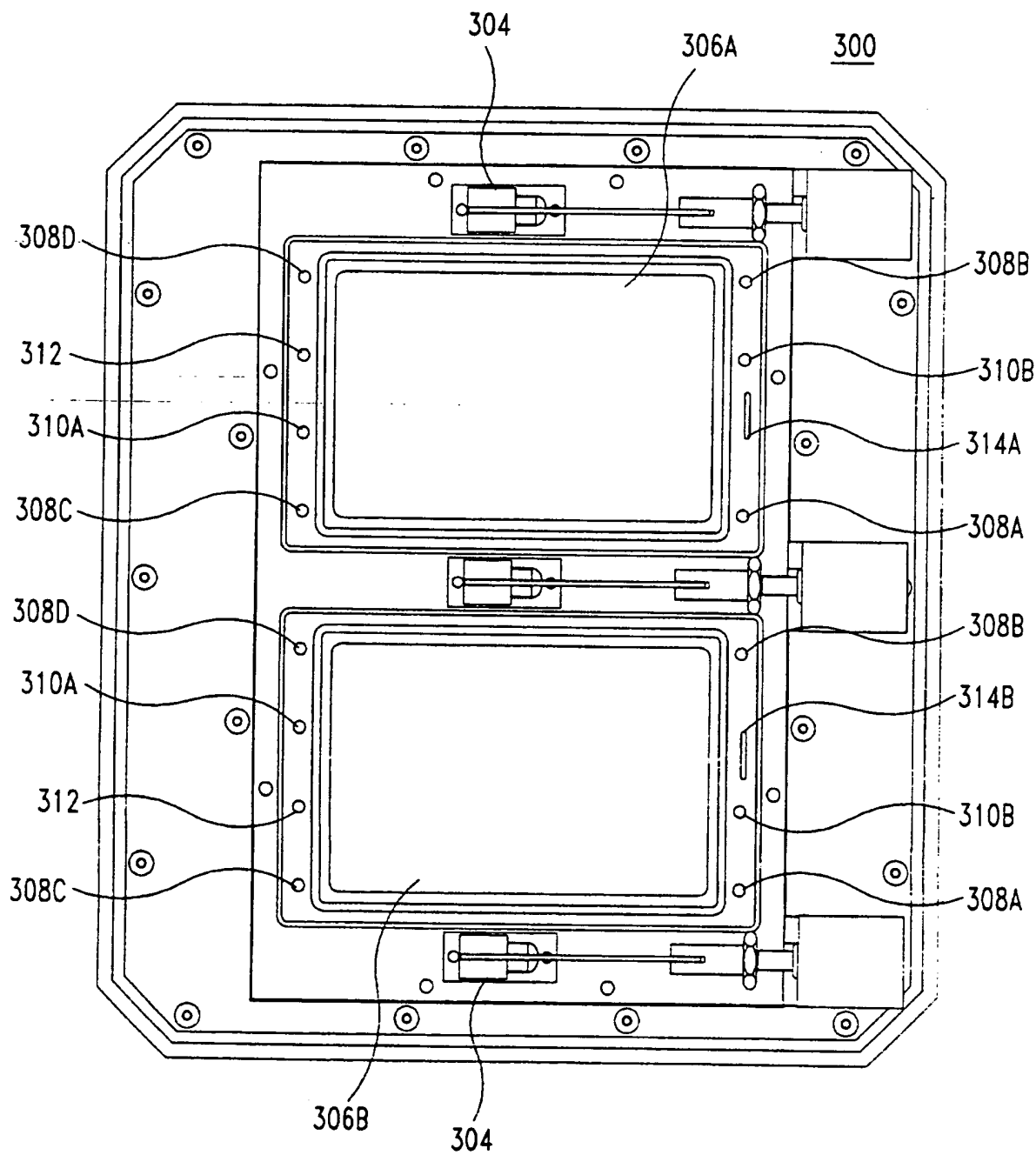


FIG. 13

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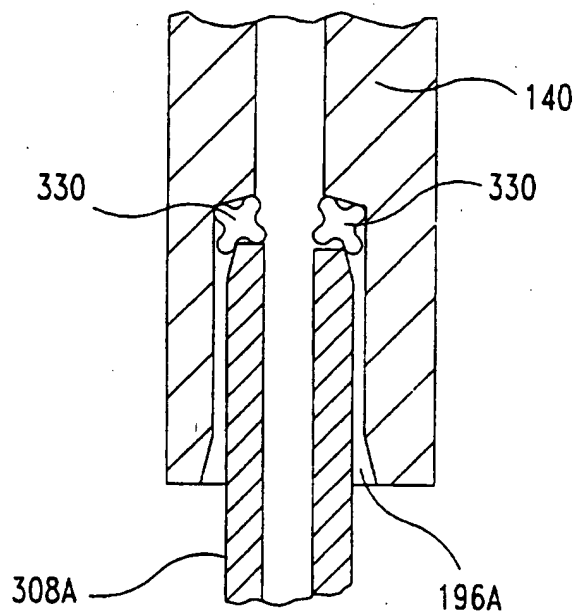


FIG. 14

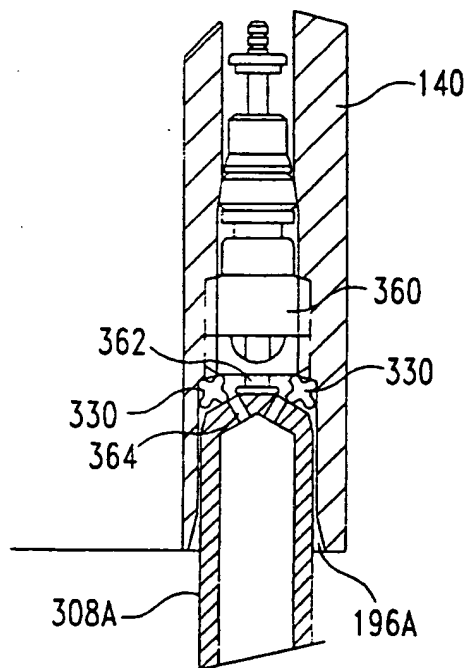


FIG. 15

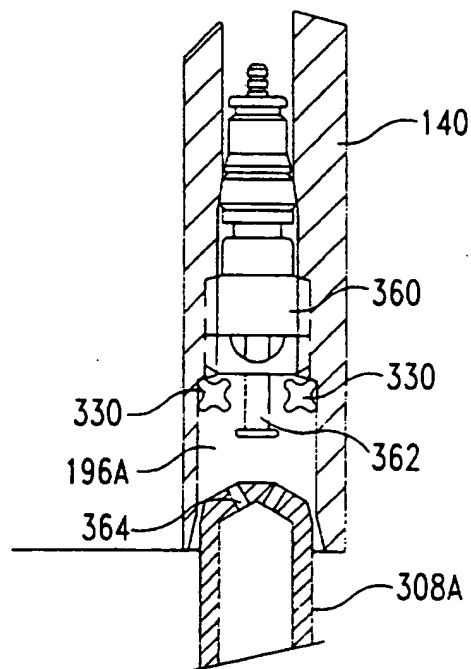


FIG. 16

## INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/05339**A. CLASSIFICATION OF SUBJECT MATTER**

IPC(6) : B01J 8/06; B01L 3/14

US CL : 422/196, 99, 102

According to International Patent Classification (IPC) or to both national classification and IPC

**B. FIELDS SEARCHED**

Minimum documentation searched (classification system followed by classification symbols)

U.S. : 422/196, 99, 102, 63, 101; 356/244, 246; 211/74

Documentation searched other than minimum documentation to the extent that such documents are included in the fields searched

Electronic data base consulted during the international search (name of data base and, where practicable, search terms used)

APS

search terms: septum, peptides, frit, filter, tube, funnel, block, synthesis, column, reactor

**C. DOCUMENTS CONSIDERED TO BE RELEVANT**

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 5,240,680 (ZUCKERMANN ET AL) 31 August 1993 (31.08.93), entire document.	1-42
Y	US, A, 4,398,382 (SUOVANIEMI ET AL) 16 August 1983 (16.08.83), entire document.	1-42
Y	US, A, 3,901,653 (JONES ET AL) 26 August 1975 (26.08.75), entire document.	6-13 and 27-36
Y, P	US, A, 5,419,874 (COASSIN ET AL) 30 May 1995 (30.05.95), entire document.	1-42
Y	US, A, 5,356,814 (CARRICO, JR. ET AL) 18 October 1994 (18.10.94), entire document.	1-42

☒ Further documents are listed in the continuation of Box C. ☐ See patent family annex.

* Special categories of cited documents:	* later document published after the international filing date or priority date and not in conflict with the application but cited to understand the principle or theory underlying the invention
*A* Document defining the general state of the art which is not considered to be of particular relevance	*X* document of particular relevance: the claimed invention cannot be considered novel or cannot be considered to involve an inventive step when the document is taken alone
*E* earlier document published on or after the international filing date	*Y* document of particular relevance: the claimed invention cannot be considered to involve an inventive step when the document is combined with one or more other such documents, such combination being obvious to a person skilled in the art
*L* Document which may throw doubts on priority claim(s) or which is cited to establish the publication date of another citation or other special reason (as specified)	*Z* document member of the same patent family
*O* document referring to an oral disclosure, use, exhibition or other means	
*P* Document published prior to the international filing date but later than the priority date claimed	

Date of the actual completion of the international search

13 JULY 1996

Date of mailing of the international search report

09 AUG 1996

Name and mailing address of the ISA/US  
Commissioner of Patents and Trademarks  
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Washington, D.C. 20531

Facsimile No. 703/305-4230

Authorized official  
ALEXANDER MARKOFF

Telephone No. 703/305-0651

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/05339

## C (Continuation). DOCUMENTS CONSIDERED TO BE RELEVANT

Category*	Citation of document, with indication, where appropriate, of the relevant passages	Relevant to claim No.
Y	US, A, 5,108,704 (BOWERS ET AL) 28 April 1992 (28.04.92), entire document.	1-121
Y	US, A, 5,039,493 (OPRANDY) 13 August 1991 (13.08.91), entire document.	43-121
Y	US, A, 5,283,039 (AYSTA) 01 February 1994 (01.02.94), entire document.	43-121
Y	US, A, 5,219,528 (CLARK) 15 June 1993 (15.06.93), entire document.	43-121
Y	US, A, 5,324,480 (SHUMATE ET AL) 23 June 1994 (23.06.94), entire document.	122-136
Y	US, A, 5,061,639 (LUNG ET AL) 29 October 1991 (29.10.91), entire document.	122-136
Y	US, A, 5,324,483 (CODY ET AL) 28 June 1994, (24.06.94), entire document.	

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/05339

## Box I Observations where certain claims were found unsearchable (Continuation of item 1 of first sheet)

This international report has not been established in respect of certain claims under Article 17(2)(a) for the following reasons:

1. ☐ Claims Nos.:  
because they relate to subject matter not required to be searched by this Authority, namely:
  
2. ☐ Claims Nos.:  
because they relate to parts of the international application that do not comply with the prescribed requirements to such an extent that no meaningful international search can be carried out, specifically:
  
3. ☐ Claims Nos.:  
because they are dependent claims and are not drafted in accordance with the second and third sentences of Rule 6.4(a).

## Box II Observations where unity of invention is lacking (Continuation of item 2 of first sheet)

This International Searching Authority found multiple inventions in this international application, as follows:

Please See Extra Sheet.

1. ☒ As all required additional search fees were timely paid by the applicant, this international search report covers all searchable claims.
2. ☐ As all searchable claims could be searched without effort justifying an additional fee, this Authority did not invite payment of any additional fee.
3. ☐ As only some of the required additional search fees were timely paid by the applicant, this international search report covers only those claims for which fees were paid, specifically claims Nos.:
  
4. ☐ No required additional search fees were timely paid by the applicant. Consequently, this international search report is restricted to the invention first mentioned in the claims; it is covered by claims Nos.:

Remarks on Protest

- ☐ The additional search fees were accompanied by the international protest.  
☒ No protest accompanied the payment of additional search fees.

# INTERNATIONAL SEARCH REPORT

International application No.  
PCT/US96/05339

## BOX II. OBSERVATIONS WHERE UNITY OF INVENTION WAS LACKING

This ISA found multiple inventions as follows:

This application contains the following inventions or groups of inventions which are not so linked as to form a single inventive concept under PCT Rule 13.1. In order for all inventions to be examined, the appropriate additional examination fees must be paid.

Group I, claims 1-121, drawn to a reaction chamber and a reaction block.

Group II, claims 122-124, drawn to a container rack.

Group III, claims 125-136, drawn to a docking station.

The inventions listed as Groups I, II and III do not relate to a single inventive concept under PCT Rule 13.1 because, under PCT Rule 13.2, they lack the same or corresponding special technical features for the following reasons:

The invention of Group I is a reaction block having a plurality of reaction chambers. The reaction block can be used for conducting chemical reactions. The reaction block does not require for its use any container rack. The reaction block does not require the special technical features of the docking station and can be used alone.

The invention of Group II is a container rack for supporting a plurality of containers. The container rack is intended to be used in a pipetting station.

The invention of Group III is a docking station. The docking station does not require a reaction block having plurality of reaction chambers. Moreover, the docking station does not have any pipetting means and does not require any container or container rack.

The inventions are not so linked by a special technical features so as to form a single general inventive concept.

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